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| (54) Title: VERTEBRATE HOMOLOGUES OF UNC-53 PROTEIN OF C. ELEGANS | | |
| (57) Abstract Vertebrate protein homologues of UNC-53 protein of C. elegans and nucleic acid sequences coding for said homologues or functional equivalents thereof are identified. The nucleic acid sequences in an appropriate vector are used to transfect or transform cells, tissues or organisms useful in identifying inhibitors or enhancers of the vertebrate homologue, or further proteins involved in the signal transduction pathway of which said vertebrate homologue is a component. Any of said inhibitors or enhancers identified can be included in a pharmaceutical composition or in the preparation of a medicament for treating conditions such as neurological diseases, acute traumatic injuries and to promote neuronal regeneration and inhibit metastasis or loss of contact inhibition. | | |

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VERTEBRATE HOMOLOGUES OF UNC-53
PROTEIN OF C. ELEGANS

The present invention relates to vertebrate
5 homologues of UNC-53 protein of C. elegans and cDNA
sequences coding for said homologues or functional
equivalents thereof. The invention also relates to
processes for identifying compounds which control cell
behaviour, compounds identified and pharmaceutical
10 compositions containing them in addition to processes
and assays for identifying disease states in which
said gene or protein is dysfunctional.

The control of cell motility, cell shape and
directionality of cell outgrowth of axones or other
15 cell outgrowths is an essential feature in the
morphogenesis and function of both unicellular and
multicellular organisms. The control of these
processes is disturbed in a variety of disease states
in which, for example, the Receptor Tyrosine kinase
20 (RTK) signal transduction pathways, or the like, or
their downstream intra-cellular pathways (which are
shared with other extra-cellular receptors, including
cell adhesion molecules like N-CAMS and integrins) are
overstimulated.

25 Some cell surface proteins and extra-cellular
molecules controlling the directionality and potential
of cell migration have been identified, although the
processes involved are not generally understood.

It is generally considered that a long-range
30 migration of a cell process (also known as a growth
cone extension) is a stepwise event, whereby prior to
and after each extension there is the formation of a
structure at the leading edge of the cell which senses
signals in the environment instructing the cell to
35 either stabilise a cell process extending in a

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preferred direction, or to cause a lamellipodium to extend a process in a given direction. Localised stabilisation of the actin cytoskeleton and association with plus end regions of microtubules is a general cell biological process underlying the choice of directional extension. Microtubule binding directing these processes has not previously been identified. The present inventors have surprisingly found that UNC-53 protein of *C. elegans* and vertebrate homologues thereof is involved in binding of microtubules and particularly of plus end regions of microtubules.

A gene from the free-living nematode *Caenorhabditis elegans* designated "unc-53" has been previously identified and cloned (Abstract, International *C. elegans* Meeting, June 1-5 1991, Madison, Wisconsin, 58, Bogaert and Goh). The present inventors previously identified UNC-53 protein as a signal transducer or signal integrator controlling the directionality of cell migration and/or cell shape in *C. elegans* (WO 96/38555). Increased UNC-53 protein activity was found to be proportional to cell process extensions in the correct direction of cell migration. The unc-53 gene was found to encode a signal transduction molecule that transduces a signal from an RTK such as, for example, via the adaptor protein SEM-5/GRB-2, to the machinery controlling directional growth cone extension or stabilisation, in a highly dosage - dependent fashion.

Genetic and experimental analysis of *C. elegans* UNC-53 mutants showed that mutations in the unc-53 gene do not affect the general ability of cells to migrate but rather affect the ability of cells to migrate under specific antero-posterior cues. Reduction of UNC-53 activity leads to loss of

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direction and reduction of growth cone extension as indicated by the directionality of random extension cycles observed in excretory canal growth cones in UNC-53 mutants.

5 The function of UNC-53 is highly sensitive to its dosage or activity. Reduction of function leads to proportional reduction of migration to the specific signal while increased expression, using transgenic expression of UNC-53 in muscle cells, leads to
10 increased directional migration. The data lead to the conclusion that UNC-53 functions as an integrator of a directional signal in the organism whereby reception of signals leads to growth cone extension in the correct direction.

15 Certain alleles of UNC-53 enhance the sex myoblast migration defect of SEM-5 C. elegans mutants in a receptor tyrosine kinase signal transduction pathway (Stern et al 1993 mol. Biol. cell, 4, 1175-1188). While the genetics suggests that UNC-53 and
20 SEM-5 cooperate to regulate sex myoblast migration, genetic experiments do not permit a conclusion that this is the result of a direct molecular interaction. The inventors previously identified a potential sem-5/GRB-2 binding site and showed in two types of
25 biochemical experiments that UNC-53 physically interacts with SEM-5. The present inventors conclude that UNC-53 encodes a signal transduction molecule that transduces extracellular signals for directional migration via the adapter protein SEM-5/GRB-2 to the
30 machinery controlling directional growth cone extension or stabilization.

 Several lines of evidence indicate that UNC-53 might act as an adapter linking extracellular signals to the actin cytoskeleton. Firstly, UNC-53 has shown
35 homology to cortical actin binding proteins and that

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it is capable of binding F-actin in vitro. In addition expression of UNC-53 in mammalian cells leads to changes in the F-actin cytoskeleton. Very low levels of UNC-53 expression increase the number of
5 filopodia and actin microspikes protruding from the cell surface. Cells expressing UNC-53 also exhibit increased neurite extension and increased cell motility. UNC-53 thus also acts as an activator of migration.

10 Considering all available data the following possible mechanisms of action of UNC-53 can be formulated.

 The choice and activation of directional growth cone extension can be accounted for by local
15 activation of UNC-53 via a SEM-5/GRB2 complex to a receptor (eg receptor tyrosine kinase signal) which reads a localized or directional signal. Changes in growth cone steering are preceded by the formation of a localized actin patch in the area of the growth cone
20 receiving the highest signal (Bentley and O'Connor et al. Curr. Op. NeuroBiol. 1994, vol 4, 43-48). UNC-53 might be directly involved in forming these actin patches through its own actin binding or cross-linking properties. Alternatively activated UNC-53
25 may (eg via its nucleotide binding domain) transduce a signal to as yet unidentified effectors. For example, activation of the small GTP-binding protein cdc42 or a related protein leads to formation of small actin patches as well as the formation of small filopodia.
30 The unc-53 pathway may be upstream of cdc42 or both signal transducers might share downstream pathways.

 The present inventors thus decided to investigate if a similar protein was present in higher organisms such as vertebrates.

35 The present inventors describe the identification

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of a family of genes in vertebrates, and particularly in man and mouse with extensive structural homology to UNC-53. The present inventors have surprisingly found that the nucleotide domains of UNC-53 from C. elegans and UNC-53 from vertebrates similarly activate motility, establishing functional equivalence. Furthermore these domains are shown to be capable of transforming NIH3T3 cells in vitro. The inventors also found changes in RNA transcripts in transformed cell lines compared to normal human tissues suggesting a role for UNC-53 in cell differentiation, morphogenesis and disease. Furthermore, in vitro assays and transgenic models are also described that identify pharmacological modulators of UNC-53 activity and assays to identify proteins interacting with UNC-53.

According to a first aspect of the present invention, there is provided a vertebrate protein homologue of UNC-53 protein of C. elegans or a functional equivalent, derivative or bioprecursor thereof, which protein homologue comprises an amino acid sequence having a statistically significant homology to the UNC-53 protein of C. elegans as illustrated in figure 2. According to the present invention a derivative should be taken to mean mutational derivatives, fusions, internal deletions, splice variants and muteins.

There is also provided according to a second aspect of the present invention a vertebrate protein homologue of UNC-53 protein of C. elegans, which protein comprises an amino acid sequence having one or more of sequence homology blocks A, B, C, D or E as illustrated in Figure 9a, or block F in Figure 12a or a sequence having a statistically significant homology therewith.

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Preferably, said vertebrate homologue is a human protein or a mouse protein.

According to a further aspect of the invention there is provided a vertebrate protein homologue of an
5 UNC-53 protein of C. elegans, which protein comprises an amino acid sequence having one or more of sequence blocks A, B, C, D, E or F which differ from those blocks of Figure 9a and Figure 12a to a significant extent only in conservative amino acid changes. In an
10 even further aspect of the invention there is provided a vertebrate protein having an amino acid sequence encoded by the nucleotide sequence from position 1 to position 6013 as illustrated in Figure 9b. There is also provided a vertebrate protein having an amino
15 acid sequence encoded by the nucleotide sequence illustrated in Figure 11d, or a functional equivalent derivative, or bioprecursor of said homologue.

According to a further aspect of the present invention there is provided a vertebrate protein
20 having an amino acid sequence corresponding to the prosite signatures as illustrated in Figure 28 for each of said homology blocks as defined above. Advantageously the prosite signatures can be used to identify a protein having a statistically significant
25 homology to the UNC-53 protein of C. elegans. (Luethy et al 1994, Protein Science, 3, 139-146).

A further aspect of the invention comprises a vertebrate homologue according to the invention comprising an amino acid sequence as shown in figure
30 9b or 11d or an amino acid sequence which differs from the amino acid sequences shown in these figures to a significant extent only in one or more conservative amino acid changes.

In a further aspect of the present invention
35 there is also provided a nucleic acid molecule, which

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is preferably DNA, and which encodes a vertebrate
homologue of UNC-53 protein of C. elegans, or a
functional equivalent derivative, fragment or
bioprecursor of said homologue according to the
5 invention. Preferably, the cDNA comprises a sequence
of nucleotides encoding an amino acid sequence as
illustrated in figures 9b or 11d or an amino acid
which differs from the sequences shown in these
figures to a significant only in one or more
10 conservative amino acid changes. Preferably the DNA is
cDNA, which cDNA comprises at least from position 1
to 6013 of the sequence shown in Figure 9b.
Alternatively the cDNA may comprise the sequence
illustrated in Figure 11d. Also provided by the
15 present invention is a nucleic acid sequence capable
of hybridising to the nucleic acid or DNA sequences
according to the invention under high stringency
conditions, which conditions are well known to those
skilled in the art.

20 The cDNA according to the invention may be
included in an expression vector which may itself be
used to transform or transfect a host cell, which cell
may be bacterial or eukaryotic in origin including
such as, for example an animal or plant cell a fungal
25 cell or an insect cell. Thus, advantageously, once
the cDNA corresponding to the genome of the vertebrate
homologue of UNC-53 of C. elegans is synthesised,
using for example, reverse transcriptase or the like,
a range of cells, tissues or organisms may be
30 transfected following incorporation of the selected
cDNA clone into an appropriate expression vector. The
expression vector according to the invention may
comprise a promoter of C. elegans or one of human
mouse or viral origin and optionally a sequence
35 encoding a reporter molecule, such as, for example,

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green fluorescent protein.

The present invention, therefore, also further comprises a transgenic cell, tissue or organism comprising a transgene capable of expressing a vertebrate homologue of UNC-53 protein of C. elegans or a functional equivalent, fragment derivative or bioprecursor of said homologue. The term "transgene capable of expressing a vertebrate homologue of UNC-53 protein of C. elegans" as used herein means a suitable nucleic acid sequence which leads to the expression of a vertebrate homologue of UNC-53 protein of C. elegans having the same function and/or activity. The transgene may include, for example, genomic nucleic acid isolated from the appropriate vertebrate or synthetic nucleic acid including cDNA. The term "transgenic organism, tissue or cell, as used herein means any suitable organism and/or part of an organism, tissue or cell, that contains exogenous nucleic acid either stably integrated in the genome or in an extrachromosomal state.

Preferably the transgenic cell comprises any of, a COS cell, HepG2 cell, MCF-7 or N4 neuroblastoma cell or a NIH3T3 cell or a colorectal or carcinoma cell or a human derived cell such as a fibroblast or the like. The transgenic organism may be an insect, a non-human animal or a plant and preferably C. elegans or a related nematode. Preferably, the transgene comprises the nucleic acid sequence encoding the vertebrate homologue or a functional fragment of said gene according to the invention as described above. The transgene preferably comprises an expression vector according to the invention.

The term "functional fragment" as used herein should be taken to mean a fragment of the gene coding for the vertebrate homologue of the UNC-53 protein of

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C. elegans or a functional equivalent or derivative or bioprecursor of said protein. For example, the gene may comprise deletions or mutations but may still encode a functional vertebrate homologue of UNC-53 protein.

Further provided by the present invention is a method of producing a mutant vertebrate non-human organism or cell having a mutation in the wild-type gene coding for the vertebrate homologue of UNC-53 protein, which mutation affects cell behaviour or the regulation of cell motility or the shape or the direction of cell migration or microtubule plus end stability or function and localisation of protein complexes located thereon, which method comprises inducing a mutation in the vertebrate homologue of UNC-53 protein in said organism or cell. These mutant organisms or cells may be used in a screen to identify the effects of compounds on these cell functions.

The vertebrate homologue of UNC-53 protein of C. elegans or the cDNA or genomic DNA encoding it or a functional equivalent, derivative, fragment or bioprecursor of said homologue, may advantageously be used as a medicament, or in the preparation of a medicament to promote neuronal regeneration, revascularisation or wound healing or the treatment of chronic neurodegenerative disorders or acute traumatic injuries or fibrotic disease or physiological events requiring the polarity of cells or epithelia. The present inventors have also found that the vertebrate homologue of UNC-53 protein plays a role in a transformed state of cells. Accordingly, the vertebrate homologue, dominant positive or negative mutants thereof, or inhibitors thereof may advantageously be used to induce or alleviate contact inhibition in a cell or in preventing cancer

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development. Typically, the above medical conditions may be treated in mammals and more preferably humans by either a homologue of UNC-53 protein or alternatively by a nucleic acid coding for such a protein. Alternatively an antisense oligonucleotide to said UNC-53 homologue may be used to prevent its expression. Examples of other nucleic acid sequences which may be used include 3' untranslated regions of mRNA which could be used to prevent transcription of the genomic sequence encoding for the vertebrate homologue of UNC-53 protein.

The vertebrate homologue of UNC-53 protein or a functional equivalent, fragment or bioprecursor of said protein may be incorporated into a pharmaceutically acceptable composition together with a suitable carrier, diluent or excipient therefor. The pharmaceutical composition may advantageously comprise, additionally or alternatively, the nucleic acid sequence according to the invention as defined above.

The present invention also provides for a method of determining whether a compound is an inhibitor or enhancer of the regulation of cell behaviour, growth, transformation, cell shape or motility or the direction of cell migration or microtubule plus end stability or function and localisation of protein complexes thereon which method comprises contacting said compound with a transgenic cell according to the invention and screening for a phenotypic change in said cell. Preferably the method can determine whether the compound comprises an inhibitor or an enhancer of the signal transduction pathway of said transgenic cell of which pathway said vertebrate homologue of UNC-53 protein, or a functional equivalent, derivative, fragment or bioprecursor of

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said vertebrate homologue is a component or whether said compound is an inhibitor or an enhancer of a parallel or redundant signal transduction pathway in said cell. The present invention also provides a method to determine that the protein in said signal transduction pathway is a vertebrate homologue of UNC-53 protein of C. elegans or a functional equivalent, fragment, derivative or bioprecursor of said vertebrate homologue.

10 Preferably, the phenotypic change to be screened comprises a change in cell shape or a change in cell motility. Where a transgenic cell is used in accordance with one embodiment of the method of the invention, an N4 neuroblastoma cell may be used and in
15 such an embodiment the phenotypic change to be screened may be the length of neurite growth or changes in filipodia outgrowth or alternatively changes in ruffling behaviour or cell adhesion or any change in microtubule cytoskeleton or any change in
20 localisation of proteins on plus end regions of microtubules or any change in cell death such as in apoptosis. In an alternative embodiment of the method of the invention, the transgenic cell may comprise an MCF-7 breast cancer cell. Typically in such an
25 embodiment the phenotypic change to be screened comprises the extent of phagokinesis or filipodia formation. In an alternative embodiment of this aspect of the invention, the transgenic cell may comprise an NIH3T3 cell. Typically in such an
30 embodiment the phenotypic change to be screened comprises loss of contact inhibition of foci formation. The method according to the invention, may also utilise a mutant cell or mutant organism according to the invention as described above, where
35 the mutant cell is capable of growing in tissue

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culture or in vivo and either of which cell or organism has a mutation in the wild-type unc-53 gene.

In accordance with the present invention, a "phenotypic change", may be any phenotype resulting from changes at any suitable point in the life cycle of the cell, tissue or organism defined above, which change can be attributed to the expression of the transgene such as for example, growth, viability, morphology, behaviour, movement, cell migration or cell process or growth cone extension of cells and includes changes in body shape, locomotion, chemotaxis, contact inhibition, mating behaviour or the like. The phenotypic change may preferably be monitored directly by visual inspection of the cell as a whole or particularly by monitoring the F-actin cytoskeleton microtubule network and plus end stability of microtubules or proteins thereon or alternatively by for example measuring indicators of viability including endogenous or transgenically introduced histochemical markers or other reporter genes, such as for example β -galactosidase or green fluorescent protein.

A compound which is identifiable by the method according to the invention as described above, as an enhancer of the processes identified above such as the regulation of cell shape or motility or the direction of cell migration may be used as a medicament, or alternatively in the preparation of a medicament, for promoting neuronal regeneration, revascularisation or wound healing, or for treatment of chronic neurodegenerative diseases or acute traumatic injuries or fibrotic disease. Examples of promoting neuronal regeneration include, for example, peripheral nerve regeneration after trauma and spinal cord trauma.

Where a compound is identified in accordance with

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the method described above as being an inhibitor of the regulation of cell shape etc., the compound may be used as a medicament, or in the preparation of a medicament, for substantially alleviating spread of disease inducing cells, such as in spread of cancer, or the like in metastasis or in alleviating loss of contact inhibition. Advantageously, any of the compounds which may have been identified as an inhibitor or an enhancer in accordance with the method as described above, may also be included in a pharmaceutical composition comprising the respective compound and a pharmaceutically acceptable carrier, diluent or excipient therefor.

The particular mechanism of action of a compound identified as either an inhibitor or an enhancer of the cell motility shape, growth or direction of cell migration or microtubule association or to the plus end region thereof is not limiting. Preferably the compound acts as an inhibitor or enhancer of a signal transduction pathway. The compound may also act on a parallel pathway or directly on the vertebrate homologue of UNC-53 protein of C. elegans. For example, the method of action of the compound may include direct interaction with the vertebrate homologue of UNC-53 protein, interaction with processes for regulating phosphorylation or dephosphorylation of the vertebrate homologue of UNC-53 or with processes regulating activity of an unc-53 gene or with processes for post-transcriptional or post-translational modification or the like.

Preferably the compound is identified by the method according to the invention as an inhibitor or an enhancer, by utilising differences of phenotype of the cell, tissue or organism, which are visible to the eye. Alternatively indicators of viability including

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endogenous or transgenically introduced histochemical markers or a reporter gene may be used.

According to a further aspect of the invention there is also provided a transgenic cell or tissue
5 culture which has been constructed to comprise a promoter sequence of a gene coding for a vertebrate homologue of UNC-53 of C. elegans or a functional equivalent, derivative fragment, or bioprecursor of said homologue operably linked to a nucleic acid
10 sequence encoding a reporter molecule. Preferably, the reporter sequence encoding the reporter molecule which comprises a detectable protein, for example one which may be monitored by eye inspection such as antibiotic resistance, β -galactosidase or a molecule
15 detectable by spectrophotometric, spectrofluorometric, luminescent or radioactive assays.

The present invention also provides a method of determining whether a compound is an inhibitor or an enhancer of transcription of a gene coding for a
20 vertebrate homologue of UNC-53 protein in C. elegans, or a functional equivalent, derivative fragment or bioprecursor of said homologue, which method comprises the steps of:

- 25 (a) contacting said compound with a transgenic cell according to the invention as described above,
- (b) monitoring the level of said reporter molecule and comparing results obtained from this monitoring step with a control comprising a
30 transgenic cell having the promoter sequence of a gene coding for a vertebrate homologue of UNC-53 protein, or a functional fragment of said homologue and the reporter molecule, in the absence of the compound.

35 In one embodiment of the method according to this

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aspect of the invention the reporter molecule may comprise messenger RNA.

5 A compound identified as an enhancer of transcription of the gene coding for the vertebrate homologue of UNC-53 protein of C. elegans or a functional equivalent, derivative or bioprecursor of said homologue may also be used as a medicament, or in the preparation of a medicament, for promoting neuronal regeneration, revascularisation or wound
10 healing, or for treatment of chronic neuro-degenerative diseases or acute traumatic injuries or fibrotic disease. Furthermore, such compounds may be included in a pharmaceutical composition including a pharmaceutically acceptable carrier, diluent or
15 excipient therefor. Any compounds identified as inhibitors of transcription may, advantageously, be used in alleviating the spread of disease inducing cells such as cancers or metastasis or loss of contact inhibition.

20 The present invention also provides a kit for determining whether a compound is an enhancer or an inhibitor of the regulation of cell growth, transformation, cell motility or shape or the direction of cell migration which kit comprises at
25 least one transgenic or mutant cell or transgenic or mutant non-human organism according to the invention as described above and a plurality of wild-type cells or one organism of the same type, or a cell line or tissue culture and means for contacting said compound
30 with said cell or organism.

Also provided by the present invention is a kit for determining whether a compound is an inhibitor or an enhancer of transcription of a gene coding for a vertebrate homologue of UNC-53 protein of C. elegans
35 or a functional equivalent, derivative or fragment

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thereof, which kit comprises at least one transgenic cell or cells according to the invention and means for contacting said compounds with said cells.

For the purposes of the present invention, the
5 term "gene coding for a vertebrate homologue of UNC-53
or a functional fragment of said homologue" includes
the nucleic acid sequence shown in Figures 9b or 11d
or a fragment thereof, including the differentially
spliced isoforms and transcriptional starts of the
10 nucleic acid sequence and which sequence encodes a
vertebrate homologue of UNC-53 protein or a functional
equivalent, derivative, fragment or bioprecursor of
the protein.

The present invention also provides methods of
15 identifying genes of vertebrates or fragments of said
genes, which encode proteins which are active in the
signal transduction pathway of which the vertebrate
homologue of UNC-53 is a component. A preferred
method comprises hybridizing to an appropriate cDNA
20 library a nucleotide sequence, as defined herein, or a
fragment thereof under appropriate conditions of
stringency in order to identify genes having
statistically significant homology with the cDNA
clones of any one of the cDNA sequences according to
25 the invention described above.

Furthermore, there is also provided by the
present invention a method of identifying a protein
which is active in the signal transduction pathway of
a cell of which a vertebrate homologue of UNC-53
30 protein of C. elegans or a functional equivalent,
fragment or bioprecursor of said vertebrate homologue
is a component. According to this aspect of the
invention, the method comprises;

(a) contacting an extract of said cell with an
35 antibody to the vertebrate homologue of UNC-53

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protein or a functional equivalent, fragment or bioprecursor of said protein,

(b) identifying the antibody/vertebrate homologue of UNC-53 complex, and

5 (c) analysing the complex to identify any protein bound to the vertebrate homologue of UNC-53 protein other than the antibody.

The vertebrate homologue of UNC-53 protein, therefore may bind regions of other proteins involved
10 in the signal transduction pathway. It is also possible to sequentially identify a whole range of proteins involved in the signal transduction pathway.

Antibodies to the vertebrate homologue of UNC-53 protein may be produced according to known techniques
15 as would be known to those skilled in the art. For example, polyclonal antibodies may be prepared by inoculating a host animal, such as a mouse, with a protein or epitope of a protein according to the invention and recovering immune serum.

20 This aspect of the invention further comprises a method of identifying a further protein or proteins which are active in the signal transduction pathway of a cell of which UNC-53 is a component which method comprises:

25 (a) forming an antibody to the first identified protein bound to the vertebrate homologue of UNC-53 protein in the method as described above,
(b) contacting a cell extract with the antibody,
(c) identifying the antibody/protein complex,
30 (d) analysing the complex to identify any further protein bound to the first protein other than the antibody, and
(e) optionally repeating steps (a) to (d) to identify further proteins in the pathway.

35 According to this aspect of the present

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invention, the antibody starts the process by binding to the vertebrate homologue of UNC-53 protein or a functional equivalent thereof in the signal transduction pathway. Any other proteins found
5 complexed to the bound antibody or UNC-53 protein can then be used to identify further interacting proteins involved in the pathway.

It may also be possible to identify proteins involved in the signal transduction pathway of a cell
10 of which the vertebrate homologue of UNC-53 or a functional equivalent derivative or bioprecursor thereof is a component by using a vertebrate homologue of UNC-53 protein of C. elegans. According to this aspect of the invention the method comprises:

- 15 (a) contacting an extract of the cell with the vertebrate homologue of UNC-53 protein of C. elegans or a functional equivalent, fragment or bioprecursor of said homologue,
- (b) identifying the vertebrate homologue of
20 UNC-53 protein/protein complex formed and
- (c) analysing the complex to identify any protein bound to the vertebrate homologue of UNC-53 protein other than the same vertebrate homologue of UNC-53 protein

25 This method can also advantageously be used to identify further proteins in a signal transduction pathway of a cell by contacting an extract of the cell used as described above, with any protein identified from step (c) above not being a vertebrate homologue
30 of UNC-53 protein and repeating steps (b) and (c).

Other methods which may be used for identifying proteins in a signal transduction pathway of a cell may comprise for example a western blot overlay method which method is well known to those skilled in the
35 art. Cell extracts are run on gels to separate out

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protein and subsequently blotted onto a nylon membrane. These membranes may then be incubated, for example in a medium containing a vertebrate homologue of UNC-53 having a label attached thereto such as a biotin or radiolabel and any protein conjugates visualised with for example a streptavidin or alkaline phosphatase conjugated antibody.

The present invention also advantageously provides a process for the preparation of binding antibodies which recognise proteins or fragments thereof involved in the rate and direction of cell migration or the control of cell growth or shape, for the above methods.

The monoclonal antibody for binding to the appropriate vertebrate homologue of UNC-53 (or its functional equivalent) may be prepared by known techniques as described by Kohler R. and Milstein C., (1975) Nature 256, 495 to 497.

Another method which may be used to identify proteins involved in the signal transduction pathway of a cell of which a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent or derivative or bioprecursor is a component involves investigating protein-protein interactions using the two-hybrid vector method. This method is well known to those skilled in the art and which was first developed in yeast by Chien et al (1991). This technique is based on functional reconstitution in vivo of a transcription factor which activates a reporter gene. More particularly the technique comprises providing an appropriate host cell with a DNA construct comprising a reporter gene under the control of a promoter regulated by a transcription factor having a DNA binding domain and an activating domain, expressing in the host cell a first hybrid DNA

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sequence encoding a first fusion of a fragment or all of a nucleic acid sequence according to the invention and either said DNA binding domain or said activating domain of the transcription factor, expressing in the host at least one second hybrid DNA sequence, such as a library or the like, encoding putative binding proteins to be investigated together with the DNA binding or activating domain of the transcription factor which is not incorporated in the first fusion; detecting any binding of the proteins to be investigated with a protein according to the invention by detecting for the presence of any reporter gene product in the host cell; optionally isolating second hybrid DNA sequences encoding the binding protein.

15 An example of such a technique utilises the GAL4 protein in yeast. GAL4 is a transcriptional activator of galactose metabolism in yeast and has a separate domain for binding to activators upstream of the galactose metabolising genes as well as a protein binding domain. Nucleotide vectors may be constructed, one of which comprises the nucleotide residues encoding the DNA binding domain of GAL4. These binding domain residues may be fused to a known protein encoding sequence, such as for example a sequence coding for the vertebrate homologue of UNC-53. The other vector comprises the residues encoding the protein binding domain of GAL4. These residues are fused to residues encoding a test protein, preferably from the signal transduction pathway of the vertebrate in question. Any interaction between the vertebrate homologue of UNC-53 protein and the protein to be tested leads to transcriptional activation of a reporter molecule in a GAL-4 transcription deficient yeast cell into which the vectors have been transformed. Preferably, a reporter

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molecule such as β -galactosidase is activated upon restoration of transcription of the yeast galactose metabolism genes. This method enables any interactions between proteins involved in the signal transduction pathway or a parallel or redundant pathway to be investigated.

Any proteins identified in the signal transduction pathway of the cell, which may be for example a mammalian cell, may also be included in a pharmaceutical composition together with a pharmaceutically acceptable carrier, diluent or excipient therefor.

The present invention also provides a process for producing a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, fragment, or derivative of the protein, which process comprises culturing the cells transformed or transfected with a cDNA expression vector having any of the cDNA sequences according to the invention as described above, and recovering the expressed vertebrate homologue of UNC-53 protein. The cell may advantageously be a bacterial, animal, insect or plant cell.

A particularly preferred process for producing a vertebrate homologue of UNC-53 protein or a functional equivalent, derivative or fragment of said homologue comprises using insect cells. Accordingly, the invention provides a process for producing a vertebrate homologue of UNC-53 protein of C. elegans or a functional equivalent, fragment, derivative or bioprecursor of the UNC-53 protein, which process comprises culturing an insect cell transfected with a recombinant Baculovirus vector, said vector comprising a nucleotide vector encoding the vertebrate homologue of UNC-53 protein or a functional equivalent, fragment

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or bioprecursor thereof downstream of the Baculovirus polyhedrin promoter and recovering the expressed protein. Advantageously, this method produces large amounts of protein for recovery. The insect cell may
5 be from for example Spodoptera frugiperda or Drosophila Melanogaster.

In accordance with the present invention, a defined nucleic acid sequence includes not only the identical nucleic acid but also any minor base
10 variations from the natural nucleic acid sequence including in particular, substitutions in bases which result in a synonymous codon (a different codon specifying the same amino acid), due to the degenerate code in conservative amino acid substitution. The
15 term "nucleic acid sequence" also includes the complimentary sequence to any single stranded sequence given which includes the definition above regarding base variations.

Furthermore, a defined protein, polypeptide or
20 amino acid sequence according to the invention, includes not only the identical amino acid sequence but also minor amino acid variations from the natural amino acid sequence including conservative amino acid replacements (a replacement by an amino acid that is
25 related in its side chains). Also included are amino acid sequences which vary from the natural amino acid but result in a polypeptide which is immunologically identical or similar to the polypeptide encoded by the naturally occurring sequence. Such polypeptides may
30 be encoded by a corresponding nucleic acid sequence.

A further aspect of the invention provides a nucleic acid sequence of at least 15 nucleotides of a nucleic acid according to the invention and preferably from 15 to 50 nucleotides.

35 These sequences may, advantageously be used as

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probes or primers to initiate replication or the like. Such nucleic acid sequences may be produced according to techniques well known in the art, such as by recombinant or synthetic means. They may also be used
5 in diagnostic kits or the like for detecting for the presence of a nucleic acid according to the invention. These tests generally comprise contacting the probe with a sample under hybridising conditions and detecting for the presence of any duplex formation
10 between the probe and any nucleic acid in the sample. Nucleic acid sequences according to the invention may also be produced using recombinant or synthetic means such as described in Sambrook et al (Molecular Cloning: A Laboratory Manual, 1989). Advantageously,
15 human allelic variants or polymorphisms of the DNA according to the invention may be identified by, for example, probing DNA libraries from a range of individuals for example from different populations. Furthermore, nucleic acids and probes according to the
20 invention may be used to sequence genomic DNA from patients using techniques well known in the art, such as the Sanger Dideoxy chain termination method, which may advantageously ascertain any predisposition of a patient to certain proliferative disorders.

25 A method of detecting whether a compound is an inhibitor or an enhancer of expression of a vertebrate homologue of UNC-53 of C. elegans, or a functional equivalent, derivative or fragment of said vertebrate homologue is also provided which method comprises
30 contacting a cell expressing said homologue with said compound and monitoring for a phenotypic change compared to a control cell which has not been contacted with said compound.

35 Preferably the cell is a transgenic cell as described above. Alternatively the cell may have

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undergone loss of contact inhibition.

The present method also provides for determining whether said compound is an inhibitor of expression of said vertebrate homologue. In one embodiment the
5 compound to be tested comprises a nucleic acid.

Preferably said nucleic acid sequence comprises an antisense DNA sequence or a mRNA sequence.

Preferably said mRNA sequence comprises 3' untranslated regions of mRNA encoding for said
10 vertebrate homologue.

Alternatively, the compound to be tested may be a protein. Preferably, said protein comprises a protein having an amino acid sequence potentially suitable for inhibiting function of said vertebrate homologue and
15 preferably comprises a protein identified by the methods as described herein.

The present invention also provides a pharmaceutical composition comprising a compound, for example an antisense nucleic acid identified according
20 to the above described method together with a pharmaceutically acceptable carrier, diluent or excipient therefor.

A nucleic acid sequence or protein identified according to this aspect of the invention may be used
25 as a medicament, or in the preparation of a medicament, for treating loss of contact inhibition or cancer which is mediated by a vertebrate homologue of UNC-53 protein or a functional equivalent, fragment, derivative or bioprecursor of said homologue.

Further provided by the invention is a nucleic
30 acid as defined above for use in preparation of a medicament for inhibiting expression of a gene coding for a vertebrate homologue of UNC-53 protein of C. elegans or a functional equivalent, derivative,
35 fragment or bioprecursor of said homologue.

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According to a further aspect of the invention there is provided a plasmid pCB201 deposited under LMBP Accession No. LMBP 3594 and a MCF-7 and a NIH/3T3 cell line transfected with plasmid pCB201 deposited
5 under LMBP Accession Nos. LMBP 1601 CB and LMBP 1603 CB respectively. Further provided by the invention is phage lambda 3b coding for Hu-UNC-53/1 and deposited under Accession No. LMBP 1604CB (or 3595). Also provided are plasmids pLM1 deposited under Accession
10 No. LMBP 3762, pLM4 (LMBP 3763), pEGFP72 (LMBP 3764) and pCB501 (LMBP 3765). Further provided is a Bac clone comprising a fragment of hu-unc-53/2 gene (LMBP 3773) and a worm strain comprising a chimeric *C.elegans* human unc53 gene deposited under LMBP
15 Accession No. LMBP-1663CB.

Further provided by the invention is an assay for detecting expression of a vertebrate homologue of UNC-53 protein of *C. elegans* in a vertebrate cell which assay comprises contacting a cell or an extract
20 thereof with an antibody to said vertebrate homologue, or a functional equivalent, derivative or bioprecursor thereof, which antibody is fused to a reporter molecule, removing any unbound antibody and monitoring for the presence of said reporter molecule.

25 Preferably the reporter molecule is an antibody conjugated to for example a flurophore such as fluorescein or alternatively to an enzyme such as strepavidin.

There is also provided a method for detecting for
30 expression of a gene coding for a vertebrate homologue of UNC-53 protein or a functional equivalent, derivative, fragment or bioprecursor thereof, which method comprises contacting a probe specific for a nucleic acid or protein sequence coding for or
35 corresponding to said vertebrate homologue or a

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functional equivalent, fragment or bioprecursor thereof with a cell extract, which probe is linked to a reporter and analysing for the presence of said reporter.

5 Preferably the probe is a complementary sequence to a region of mRNA transcribed from said gene encoding said vertebrate homologue of UNC-53 protein or a functional equivalent, derivative or bioprecursor therefor.

10 Preferably the complimentary sequence is a 3' or 5' untranslated region of said mRNA. Preferably said reporter may be a dig label, a fluorophore, a hapten or a radiolabel.

 Alternatively said probe comprises an antibody
15 specific for said vertebrate homologue of said UNC-53 protein or a functional equivalent, derivative, fragment or bioprecursor therefor.

 Preferably the reporter is an antibody conjugated to for example a fluorophore such as fluorscein or
20 alternatively an enzyme such as streptavidin.

 As described above UNC-53 protein of C.elegans has been found to localise to microtubule and particularly to microtubule (+) ends. Therefore, there is provided by a further aspect of the present
25 invention a method of determining whether a compound is an inhibitor or an enhancer of association of UNC-53 or a vertebrate homologue thereof according to any of claims to 1 to 9 to microtubules or plus end regions thereof, which method comprises (a) contacting
30 said compound with a transgenic cell, tissue or organism expressing UNC-53 protein or said vertebrate homologue and which protein is operably linked to a reporter molecule (b) screening for the localisation of said reporter molecule as compared to a cell
35 according to step (a) which has not been contacted

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with said compound.

A compound identifiable by the above method also forms part of the present invention. Such a compound identified as an inhibitor of localisation or
5 association of UNC-53 or said vertebrate homologue with microtubules or the plus end region thereof may be used in alleviating the spread of disease inducing cells or metastasis or loss of contact inhibition. Further a compound identified as an enhancer of
10 association of UNC-53 or said vertebrate homologue with microtubules or the plus end region thereof may be used in for example promoting neuronal regeneration, revascularisation or wound healing, or for treating chronic neurodegenerative diseases or
15 acute traumatic injuries or fibrotic disease. These compounds may then be included in a pharmaceutical composition, together with a pharmaceutically acceptable carrier, diluent or excipient therefor.

Also provided by the present invention is a kit
20 for determining whether a compound is an inhibitor or an enhancer of association of UNC-53 or a vertebrate homologue thereof according to the invention with microtubules or the plus end regions thereof, which kit comprises at least one transgenic cell expressing
25 UNC-53 and a reporter molecule or a host or transgenic cell according to the invention and at least one cell of the same cell type for use as a control and means for contacting said compound with one of said at least one transgenic cells. Compounds identified as
30 inhibitors or enhancers or microtubule association described above may advantageously be included in a composition and linked to unc-53 protein of C.elegans or a vertebrate homologue thereof according to the invention to target the compounds to the microtubules
35 or the plus end regions thereof. Such a composition

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may also comprise, for example, a suitable transfecting or transformation agent.

According to a further aspect of the invention there is provided a method of targeting a protein to a cell microtubule or the plus end region thereof, which method comprises introducing into a host cell, tissue or organism a transgene comprising a sequence capable of expressing UNC-53 or a vertebrate homologue thereof according to the invention, which sequence is operably linked to a sequence encoding said protein to be targeted such that a chimeric protein is expressed and which results in targeting said protein to said microtubule or a plus end region thereof. An even further aspect of the invention comprises a method of identifying a molecule which covalently modifies UNC-53 or a vertebrate homologue thereof according to the invention, which method comprises a) contacting either an extract from a cell or cells expressing UNC-53 or said vertebrate homologue or a mixture of enzymes comprising candidate UNC-53 modifying enzymes in the presence of an indicator of covalent modification of a protein, b) identifying any covalently modified UNC-53 protein from step a) and c) identifying said molecule involved in said modification step. Such an indicator may be ³²P.

Further provided by the invention is a method of identifying a compound which alleviates or enhances the toxicity of UNC-53 or a vertebrate homologue thereof according to the invention, or which alleviates or enhances apoptosis. The method of the former comprises contacting said compound with a transgenic cell, tissue or organism according to the invention and monitoring for the presence of said reporter molecule adjacent said microtubules or the plus end regions thereof. In the case of apoptosis the

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method comprises monitoring the effect of the compound on cell death.

The invention may be more clearly understood from the following examples which are only exemplary, with
5 reference to the accompanying drawings wherein

Figure 1 illustrates the sequence of plasmid pTB72 which codes for the full length UNC-53 protein in C. elegans, deposited under LMBP Accession No. 3486.

10 Figure 2 illustrates the full-length UNC-53 protein from C. elegans.

Figure 3 is a Tblastn search of the EST division of Genbank with the ORF of the longest known Ce-UNC-53 cDNA. tb3-M5, reveals two EST's with homology to a predicted coiled-coil region in Ce-UNC-53.

15 Figure 4 illustrates a search of the Genbank databases with part of the nucleotide binding domain of Ce-UNC-53. It does not identify statistically significant proteins except for the C. elegans cosmid containing Ce-unc-53.

20 Figure 5 illustrates a three frame translation of EST gb:R41071.

Regions of homology with Ce-Unc-53 in two different frames are underlined. The spacing between the blocks of homology is of similar size to that in
25 Ce-UNC-53. Subsequent re-cloning and re-sequencing of this region in man identified multiple sequencing errors gb:R41071, and identified an ORF which is more homologous to and co-linear with Ce-UNC-53 (see alignment in fig. 12).

30 Figure 6 is a BLASTN search of the EST division of Genbank with Hu-unc-53/1 cDNA cosmid 3b.

Figure 7 is a TBLASTN search of the Genbank sequence database with the 961 amino acid ORF of cDNA 3b of hu-UNC-53/1 : hu-UNC-53/1 forms a unique pair
35 with Ce-UNC-53 (cosmid F45E10) compared to the rest of

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the database.

Figure 8 is a diagram illustrating the length and overlap and tissue source of the different cDNA clones of the 3' end of Hu-UNC-53/1 isolated in this work.

5 Figure 8a. is a diagram illustrating the further sequence of the Hu-UNC-53/1 and overlap of constructs to obtain the further sequence.

10 Figure 8b is a diagram illustrating the 3' end of Hu-UNC53/1 and the EST clones present in the database.

Figure 9a is an annotated sequence listing of clone 3b of hu-UNC-53/1 including the EcoR1 polylinker GAATTC. The predicted Open Reading Frame of Hu-UNC-53/1 is listed below the sequence. Blocks A B C D and
15 E which are similiar to Ce-UNC-53/1, a region which is different between Hu-UNC-53/1 and Hu-UNC-53/2 and the 3' untranslated leader sequences are marked with arrows and labelled.

Figure 9b is an annotated sequence listing of
20 Hu-UNC-53/1 available at this moment. The predicted Open Reading Frames of Hu-UNC-53/1, pLM1, pLM3, pLM4, pCB251, pLM5 and pCB201, the homology blocks A,B,C,D and E, the position of a region which is different between Hu-UNC-53/1 and Hu-UNC-53/2, the position of
25 phh14-3, pCB212, pCB210-14, phh3b, phh15, the position of the reverse primers HU53rv1, HU53rv2, HU53rv3 and HU53rv4, the position of peptides B72628 (=28/1), B72627, B72626 and B72625 are listed below the sequence.

30 Figure 10 is an annotated sequence listing of the insert of clone gbAA049124 (EST479167) of mu-UNC-53/1. The open reading frame and 3' untranslated sequence is marked with an arrow.

35 Figure 11a is an annotated sequence listing of the insert of clone gbH09036 (EST46037) of Hu-UNC-

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53/2.

Figure 11b is a novel DNA sequence of HU-UNC-53/2 extended by RT-PCR. This DNA sequence is not present in EST-46037 and extends the ORF beyond position 1109 of Figure 11a to an ORF from position 18 to 1793.

Figure 11c summarises how the 3' and 5' extensions of hu-unc-53/2 were made.

Figure 11d compiles the sequence of hu-unc-53/2. The boxed sequences are the primer sequences used for the respective extension steps described in the experimental methods section.

Figure 11e illustrates the sequences of the extensions summarised in figure 11c.

Figure 11f illustrates the sequence information illustrating four alternative Start sites observed for hu-unc-53/2.

Figure 12 is an illustration of a Tblastn search of the EST division of Genbank with 680aa starting at the C-terminus of the alpha actinin domain of hu-unc-53/2.

Figure 12a is an illustration of an amino acid alignment of the available sequence of C.elegans unc-53 and hu-unc-53/1 and hu-unc-53/2.

12b. is an illustration of similarity plots for Ce-unc-53 and hu-unc-53/1 (top) and for hu-unc-53/1 and hu-unc-53/2.

Figure 13 is an annotated sequence listing of expression vector pCB201 containing homology block E from Hu-UNC-53/1 cloned in a pcDNA3.1-HIS expression vector. The HIS and T7-tags, PCR primer used to modify hu-UNC-53/1 and ORF are marked.

Figure 14 is a diagram showing the alignment of the homologous regions of hu-UNC-53/1 and mu-UNC-53/1.

Figure 15 is an annotated sequence listing of expression vector pCDU3 containing part of Ce-UNC-53/1

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cloned in expression vector pcDNA3.1. The upper ORF starts in the vector polylinker. The lower ORF starts at the first Methionine and is part of Ce-UNC-53/1.

5 Figure 16 is an annotated sequence listing of expression vector pCDU4 containing part of Ce-UNC-53/1 cloned in expression vector pcDNA3.1. The upper ORF starts in the vector polylinker. The lower ORF starts at the first Methionine and is part of Ce-UNC-53/1.

10 Figure 17 is an annotated sequence listing of expression vector pCDU2 containing part of Ce-UNC-53/1 cloned in expression vector pcDNA3.1. The upper ORF starts in the vector polylinker. The lower ORF starts at the first Methionine and is part of Ce-UNC-53/1.

15 Figure 18 illustrates MCF-7 cells transfected with pCB201 (upper) compared to mock transfected MCF-7 cells (phase contrast image). The control cells are spread out on the tissue culture plastic and exhibiting few filopodia outgrowths. The transfected cells appear smaller because they are slightly rounded up and have multiple filopodia outgrowths per cell (arrowheads).

Figure 19 is a phase contrast image of MCF-7 cells, transfected with pcDNA3.1 (19a), pCDU4 (19b), pCDU3 (19c), pCDU2 (19d) and pTB72 (19e).

25 Figure 20 is an F-actin pattern (visualized with TRITC-Phalloidin) of MCF-7 cells transfected with pcDNA3.LacZ (top panel) and with pCB201 (middle and lower panel).

30 Figure 21 is an F-actin pattern Phalloidin (visualised with TRITC-Phalloidin) of MCF-7 cells transfected with pcDNA3.1 (21a), pCDU4 (21b), pCDU3(21c), pCDU2 (21d) and pTB72 (21e).

35 Figure 22 is a phase contrast image of N4 neuroblastoma cells transfected with pcDNA3.1 (22a), pCDU4 (22b), pCDU3 (22c), pCDU2 (22d) and pTB72 (22e).

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Figure 23 is an F-actin pattern Phalloidin (visualised with TRITC-Phalloidin) of N4 neuroblastoma cells transfected with pcDNA3.1 (23a), pCDU4 (23b), pCDU3 (23c), pCDU2 (23d) and pTB72 (23e).

5 Figure 24 illustrates phase contrast images of small (top), medium (middle) and large foci (bottom) induced in a monolayer of NIH3T3 cells by transfection with pCB201.

10 Figure 25(c) illustrates human metaphase chromosomes probed with a probe 1p34 and figures 25a and 25b indicating the chromosomal location of hu-UNC-53/1 in 1q31. Essentially the same techniques were used to assign the gene hu-unc-53/2 to chromosome locus 11p15 (25d and e) as illustrated in micrograph 15 25f.

The ideograms 25a and 25d are from the International System for Human Cytogenic Nomenclature 1985. The ideograms 25b and 25e in which the relative band positions and arm ratios were derived from actual 20 chromosome measurements is from Cytogenet Cell Genet 65:206-219 (1994).

Figure 26 is an expression pattern of HU-Unc53/1 and HU-Unc532 in normal human tissues and cancer cell lines.

25 Figure 27 is a sequence map of Plasmid pNP3.

Figure 28 is an exemplary list of prosite signatures which can be used to define and identify vertebrate homologues of UNC-53.

30 Figure 29 is a annotated sequence map of plasmid pEGFPsac. The GFP-C.elegans unc53sac fusion protein, and the C.elegans unc53 sac fragment are indicated.

Figure 30 is a sequence map of plasmid pEGFP72. The GFP-C.elegans unc53 fusion protein and the C.elegans unc53 fragment are indicated.

35 Figure 31 is an annotated sequence map of plasmid

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pEGFPsma. The GFP-C.elegans unc53sma fusion protein, and the C.e.unc53 sma fragment are indicated.

Figure 32 is an annotated sequence map of plasmid pEGFPec1. The GFP-C.elegans unc53ec1 fusion protein, and the C.elegans unc53 ec1 fragment are indicated.

Figure 33 is an annotated sequence map of plasmid pEGFPxba. The GFP-C.elegans unc53xba fusion protein, and the C.elegans unc53 xba fragment are indicated.

Figure 34 is an annotated sequence map of plasmid pLM4. Open reading frames of the hui-unc53/1 and GFP are indicated.

Figure 35 is a sequence map of plasmid pNP8.

Figure 36 is an illustration of microtubule association of C.elegans Unc53, shown in HepG2 cells, transiently transfected with pTB72, expressing C.elegans Unc53. panel A: microtubule staining of HepG2 cells, using YL1/2 panel B: C.elegans Unc53 staining, using rab4.

Figure 37 is an illustration of microtubule plus-end association in human cell lines transiently transfected with pTB72, expressing C.e.Unc53. C.elegans Unc53 was stained with mab-16-48. Panel C: COS cells showing microtubule association panel B: MCF7 cells showing microtubule plus-end association panel A: HepG2 cells showing microtubule plus-end association.

Figure 38 is an illustration of microtubule association in N4 cells transiently transfected with pEGFP72, expressing the GFP-C.elegans Unc53 fusion protein. GFP fluorescence was observed in living cells. Panel A: microtubule association of the GFP-C.elegans unc53 fusion protein panel B: microtubule plus-end association of the GFP-C.elegans unc53 fusion protein.

Figure 39 is an illustration of microtubule

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association in N4 cells transiently transfected with pEGFP72, expressing the GFP-C.elegans Unc-53 fusion protein. Microtubules were stained with YL1/2 after paraformaldehyde fixation. Panel A: Microtubule association of the GFP-C.elegans unc53 fusion protein. Panel B: tubuline staining. Panel C: panel A plus panel B: co-localisation of the GFP-C.elegans unc-53 fusion protein and Tubuline can be seen as yellow.

Figure 40 is an illustration of microtubule association in N4 cells, transiently transfected with pEGFPsma, expressing the GFP-C.elegans unc53sma fusion protein. Panel A: Microtubule association of the GFP-C.elegans unc53sma fusion product. Panel B: Centriole association of GFP-C.elegans unc53sma fusion product when expressed at low levels.

Figure 41 is an illustration of microtubule association in N4 cells, transiently transfected with pEGFPec1, expressing the GFP-C.elegans unc53ec1 fusion protein. Panel A: Microtubule association of the GFP-C.elegans unc53ec1 fusion product. Panel B: Centriole association of GFP-C.elegans unc53ec1 fusion product when expressed at low levels.

Figure 42(a)/Figure 42(b) are illustrations of fluorescence of GFP in N4 cells transiently transfected with pEFPxba and pEFGPsac respectively.

Figure 43 is an illustration of microtubule association of in N4 cells transiently transfected with pLM4 expressing GFP-Hu-UNC53/1 fusion protein. Panel A: microtubule association of GFP-HU-UNC53/1 fusion protein. Panel B: microtubule plus-end association of GFP-Hu-UNC53/1 fusion protein. Panel C: microtubule association of GFP-Hu-UNC53/1 in dividing cells (end of division).

Figure 44 is an illustration of the sequence of Plasmid pNP9.

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Figure 45 is an illustration of immuno
fluorescence in melanoma G361 cells stained with sera
28.1. Panel A: Microtubule plus-end association of
Hu-UNC53/1. Panel B: microtubule plus-end association
of hu1-Unc53 in growth cone extensions.

Figure 46 is an illustration of GFP fluorescence
and immunofluorescence in N4 cells transiently
transfected with pLM4, and stained with sera 28.1.
Panel A: Fluorescence of GFP-Hu-UNC53/1 fusion
protein. Panel B: Immunofluorescence of serum 28.1.

Figure 47 is an overview of the microtubule (+)
end, the microtubule and f-actin cytoskeleton binding
properties of different constructs

Figure 50 is an illustration of rescue of lateral
ALN neurons in mutant *unc-53*.

Dorsal view of the ALN neurones axones visualise
in GFP fluorescence with the transgene *pA/GFP* in the
posterior of an adult, (c) cellular body.
a) wild type, anterior axon (aa) migrates in a
straight line along the body until reaching the head,
on the dorsal sublateral cord, posterior axon (ap)
migrates into the tail;
b) *unc-53(n152)*, anterior axons are the shorter, stop
ahead of the vulva region and form numerous collateral
branches towards the dorsal cord;
c) *unc-53(n152), pA/unc-53* anterior axons no longer
form branches and migrate in a straight line into the
head, as in the wild type at a).
scale bar 10 μ m.

Figure 51a : is an illustration of chimeric
fusion between *C. elegans* and human 1 homologue of the
unc-53 gene. The region of the putative nucleotide
binding domain (NTP) is replaced in the *C. elegans*
cDNA by the same region of the human homologue 1 of
unc-53 (H1). The cDNA is under the promotor region A

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(pA) of unc-53, which raise expression in the ALN lateral neurons.

Figure 51b : is an illustration of the chimeric minigene nematode/human pA/unc-53-H1 partially rescue the defect in the longitudinal migration of the lateral neurons ALN and PLN. The four strains compared are : wt; unc-53(n152); unc53(n152),pA/unc-53; unc-53(n152),pA/unc-53-H1. The observed phenotypes are put in three classes :

$\frac{1}{2}$ sauvage η , the axon is straight, unbranched, and migrates until the head; $\frac{1}{2}$ vulve η , the axon is straight, unbranched, and stops in the vulva region; $\frac{1}{2}$ mutant η , the axon is short, never joints the vulva region and made a lot of collateral branches. Numbers are in percentage. The number of observed axons are noted in the last column. The chimeric fusion between the C. elegans gene and human homolog (unc-53-H1) partially rescues the mutant phenotype. The chimeric gene was maded by replacing the putative nucleotide binding region (NTP)of the nematode cDNA by the same region of the human homolog 1 (H1).

Figure 52 is an illustration of the sequence for plasmid pLM5.

Figure 53 is an illustration of the sequence for plasmid pLM6.

Figure 54 is an illustration of the sequence for plasmid pLM1.

Figure 55 is a sequence map of plasmid pCB251.

Figure 56 is a sequence map of plasmid pNP10.

Figure 57 is a sequence map of plasmid pCB501.

Figure 58 is a sequence map of plasmid pTB115.

Figure 59 is a sequence map of plasmid pPD95.75.

Figure 60 is a sequence map of clone X16.

Figure 61 is a sequence map of plasmid pLM3

DEPOSITED MATERIALS

| | Deposit | Date | Acc. Nr |
|----|-------------------------------------------------------|------------------|-------------|
| | pCB201 plasmid DNA in E. coli | 3 December 1996 | LMBP 3594 |
| 5 | Lambda clone 3B encoding hu-unc-53/1 | 3 December 1996 | LMBP 3595 |
| | MCF-7 clone z4 (mock) | 3 December 1996 | LMBP 1600CB |
| | MCF-7 clone (pCB201) | 3 December 1996 | LMBP 1601CB |
| | NIH-3T3 mock | 3 December 1996 | LMBP 1602CB |
| | NIH-3T3 pCB201 | 3 December 1996 | LMBP 1603CB |
| 10 | pLM1 | 13 November 1997 | LMBP 3762 |
| | pLM4 | 13 November 1997 | LMBP 3763 |
| | pEGFP72 | 13 November 1997 | LMBP 3764 |
| | pCB501 | 13 November 1997 | LMBP 3765 |
| | BAC clone comprising fragment of hu-unc53/2 gene | 15 November 1997 | LMBP 3773 |
| 15 | Worm strain with chimeric C.elegans/human unc-53 gene | 15 November 1997 | LMBP-1663CB |

20

The above plasmids and cell lines were deposited at the Belgian Coordinated Collections of Microorganisms (BCCM) at laboratorium voor moleculaire biologie - plasmidencollective (LMBP) B9000, GENT, Belgium, in accordance with the provisions of the Budapest Treaty of 28 April 1977.

25

The present invention will now be described with reference to the following examples which are not limiting.

30

Identification of a human homologue of the UNC-53 protein of C.elegans.

Extensive searches with the ce-UNC-53 sequence (Figures 1 and 2) against the public domain databases

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(EST, Genbank, EMBL, Swissprot and PIR) revealed no statistically significant homologies (a smallest sum probability (ssp) of $10 e - 8$ is generally accepted to be significant at amino acid level). Two ESTs

5 gbH09036 (ssp = $1.1 e - 5$) a Homo sapiens cDNA clone and gbAA049124 (ssp= $8.6-5$) a mouse cDNA clone showed homology to a "coiled coil" region a common motif in the contributing to protein secondary structure. (figure 3)

10 All other candidate scores were are at background level (ssp >0.21). Careful examination of weak candidate ESTs identified EST gb:R41071 from Homo sapiens, which had obtained a low score of 53 and a non-significant probability score of 0.33 (Fig. 4).
15 The inventors surprisingly discovered potentially significant homology with the Ce-UNC-53 nucleotide binding domain, provided multiple frameshifts and sequence errors were hypothesized.

The inventors amplified, cloned and sequenced
20 part of gb:R41071 from human heart and human lung cDNA and from human genomic DNA and discovered that clone gb:R41071 had up to ten 10 different mistakes in the region checked. 5 extra nucleotides were scattered along its sequence and two nucleotide substitutions
25 were identified, and gb:R41071 lacked three nucleotides present in our clone (Fig. 5). The novel sequence obtained was two nucleotides shorter and showed the two UNC-53-homologous regions in frame. The genomic fragment obtained is larger (700 bp total
30 length) than the corresponding cDNA clones indicating the presence of an interverting sequence of around 500 bp in nucleotide 162 of this fragment. The amplified cDNA fragment which was cloned to vector PCRII (Intvitrogen) and named pCR231 and was used as a probe
35 to screen cDNA libraries.

- 40 -

The conceptual translation of the clones we obtained by PCR were screened using blast and tblastn against all known protein and DNA sequences in the database. The only clone which came up with statistically significant similarity was Ce-UNC-53 (Fig.6). This human clone and Ce-UNC-53 thus form a unique homologous pair compared to the rest of the known sequences, indicating the statistical relevance and novelty of our discovery. We designate this human gene as hu-UNC-53/1. Human heart and a human colorectal adenocarcinoma cDNA libraries were probed with pCR231 probe to identify longer cDNA clones. The clones overlap giving a linear sequence of 3706 bp (Fig 8 and 26). This sequence shows an 959 amino acid open reading frame from the beginning of the clone. The absence of a 5' untranslated region suggests that the mRNA will extend 5'.

Sequence alignment searches of the public domain databases with the DNA sequence of hu-UNC-53/1 and its' conceptual translation identified a series of ESTs most of which correspond to the 5' UTR region. (Figures 7 and 8). Surprisingly, hu-UNC-53/1 identified also the cDNA clones gbH09036 and gbAA049124 homologous to the predicted coiled coil region in Ce-UNC-53 hu-UNC-53/1, and furthermore identified a third weakly homologous EST gbR21023. The inserts of gbH09036, gbAA049124 and gbR21023 were obtained from the Merck consortium and sequenced.

gbAA049124 is >95% identical to Hu-UNC-53/1 over 604 available amino acids (fig. 10) and is the mouse orthologue of Hu-UNC-53/1. The insert in gbH09036 is clearly homologous to hu-UNC-53/1 but derived from a different locus. We therefore name the gene identified by gbAA049124 Mu-UNC-53/1 and the gene identified by gbH09036 Hu-UNC-53/2. (Figure 11).

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5 domains of high similarity mark the *unc-53* gene family

5 Ce-UNC-53 and the here-identified vertebrate
homologues form a unique novel protein family, that is
distant from the remainder of the proteins in the
public domain. Alignment of the predicted open
reading frames shows that Hu-UNC-53/1 and Hu-UNC-53/2
10 are equidistant from Ce-UNC-53. The highest homology
is found in the carboxyterminal amino acids of Ce-UNC-
53 region. The presence of a conserved GXXGKS/T box
suggests a nucleotide binding function. However, this
domain as a whole does not belong to a class of known
15 nucleotide binding proteins.

 The similarity amongst the presently known
sequence of the UNC-53 family of proteins is highest
in 5 blocks over most of the available sequence (959
amino-acids) and a firther block identified in Figure
20 12a. These blocks can be assigned signature sequences
as displayed in figure 28 or can be assigned weight
matrices based on the alignment between the different
family members. By using truncated constructs of Ce-
unc-53, the functional relevance of these domains has
25 been addressed.

**Hu-UNC53/1 and Hu-UNC-53/2 are complex
transcription units.**

30 1. A cancer cell line RNA blots probed with HU-
Unc53/1.

 A Northern blot of poly-A+RNA from several
cancer cell lines (Melanoma G361, Lung Cancer A549,
Colorectal Adenocarcinoma SW480, Burkitt Lymphoma
35 DRajii, Leukemia Molt4, Lymphoblastic Leukemia K562,

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HeLa S3 and Promyelocytic Leukemia HL60) was probed using the whole insert of pHH3b. No or weak expression was detected in the Burkitt Lymphoma DRajii, the Leukemia Molt4 and the Promyelocytic Leukemia HL60 cell lines. Five different transcripts are detected in the remaining cancer cell lines: transcripts 1 and 2 are larger than 9.5kb, transcripts 3 and 4 are 6 to 7 kb and the fifth transcript is around 6 kb. Transcripts 1 and 2 are present in all experssing cell lines. Transcripts 3 and 4 are restricted to Melanoma G361, Lung Cancer A549 and Colorectal Adenocarcinoma SW480 and are the predominant transcripts in Melanoma G361 and Colorectal Adenocarcinoma SW480. Transcript 5 is restricted to Lymphoblastic Leukemia K562 and HeLa S3 and is predominant in HeLa S3.

2. Cancer cell lines RNA blots probed with HU-UNC-53/2.

A similar set of cancer cell line Northern blots were probed with a 652bp fragment of EST46037 amplified by using the primers 5'-aggagatgaagctgacagatatcc and 5'-aaacaccagtgtgagtc. HU-UNC-53/2 is expressed in Melanoma G361, Colorectal Adenocarcinoma SW480, Lymphoblastic Leukemia K562 and HeLa S3. No expression was detected in Lung Cancer A549, Burkitt Lymphoma DRajii, Leukemia Molt4 and promyelocytic leukemia HL60. Interestingly only 2 transcript sizes were detected of around 7 kb expressed in Lymphoblastic Leukemia K562 and HeLa S3 and a transcript of >9.5 kb in Melanoma G361 and Colorectal Adenocarcinoma SW480.

3. Normal Human tissue probed with HU-Unc53/1.

A Northern blot of poly-A+RNA from normal

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human tissue was probed using the whole insert of phage HH3b. Expression levels are low in all tissues with the highest level in heart and placenta, several fold lower levels in brain and testis, even lower levels in skeletal muscle, pancreas, thymus, colon, small intestine, ovary and prostate. Expression in peripheral blood leukocyte, lung, liver, kidney, spleen is barely detectable.

4. Normal Human tissue probed with Hu-unc53/2.

A similar set of blots were probed with a 652bp fragment of EST46037 amplified by using the primers 5'-aggagatgaagctgacagatatcc and 5'-aaacaccagtgagtcc. Expression levels are low in all tissues with the highest level in kidney, lower levels in heart, placenta, lung, skeletal muscle and pancreas. Expression is barely detectable in brain and liver.

The hu-UNC53/1 and hu-UNC-53/2 homologues are clearly highly regulated genes, showing a strong tissue specificity and, probably, additional mechanisms of regulation (ie differential splicing of different promoters). The different proteins derived from RNA's identified by probe hh15 presumably share the carboxyterminal nucleotide binding domain. Ce-UNC-53 was shown to be a complex genetic locus and complex transcription unit. The different transcripts are thought to be a mechanism to assure the necessary specificity and functional diversity of this signal transduction pathway, with respect to different signals and receptors, different tissues and different directions of migration. The occurrence of a new transcript or the observed changes in expression levels in the cancer cell line blot suggests a role

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for hu-UNC-53/1 and hu-UNC-53/2 in the establishment or maintenance of the transformed state of those cells.

5 **Phenotypic changes in cells transfected with the Nucleotide Binding Domain of Ce-UNC-53/1 and Hu-UNC-53/1**

10 Ectopic expression of full length Ce-UNC-53 in *C. elegans*, murine neuroblastoma cells or human MCF-7 breast-carcinoma cells, has been found to lead to increased filopodia outgrowth and increased motility (unpublished). The structure of Ce-UNC-53 protein is reminiscent of that of large kinases or dynamin where
15 a catalytic domain is positively or negatively regulated by domains that interface with signal transduction pathways for example (by GRB2 binding, phosphorylation or the like). The inventors therefore decided to test whether the nucleotide domain by
20 itself is capable of inducing the observed changes in the microfilament cytoskeleton and motile or ruffling behaviour.

 cDNA fragments coding for the nucleotide binding domains of Ce-UNC-53 and Hu-UNC-53/1 were cloned in
25 mammalian expression vectors with the CMV promoter (see experimental procedures).

 To be able to detect expression from pCB201 (Fig. 13), an N-terminal his and a T7 epitope tag were fused in frame with the hu-UNC-53/1 cDNA hh15. pCDU3
30 contains a larger fragment of Ce-UNC-53 and starts just before the conserved "VIELKIEL" domain (Fig. 12).

 The empty pCDNA3 vector or pCDNA3.1-His-LacZ, a mammalian expression vector for *E. coli* Beta-galactosidase, was used as a control vector (mock
35 transfection). The differences between mock and

- 45 -

transfected N4 and MCF-7 clones were analysed using phase-contrast and Nomarski microscopy coupled with time lapse analysis, phagokinesis and immunocytochemical characterisation of the F-actin.

5

Phenotypic changes in mouse N4 neuroblastoma cells

10 N4 neuroblastoma cells were stably transfected with control construct pCDNA3.1 and the C. elegans UNC-53 constructs pTB72, pCDU2, pCDU3 and pCDU4. The population of clones transfected with the empty expression vector were homogeneous and similar to wild type N4 cells. In contrast thereto, 1/4 to 50% of the clones transfected with pTB72, pCDU2, pCDU3 and pCDU4 (see experimental procedures and Figs. 1,17,15 and 16 respectively) had distinct phenotypes:

1. Wild type or N4 cells transfected with pCDNA3, designated as mock transfection show a central cell body, with extensions, designated as neurite outgrowths. Less than 5% of the population have lamellae. When present, they are generally situated on the cell body and on the opposite site of the neurite extensions (figure 22a). The lamellae show a radial actin spike pattern. Limited branching of the actin fibres is observed in wild type or pCDNA3 transfected N4 cells. Side branches are smaller and can be clearly distinguished from the main actin branch (figure 23a).

2. N4 cells, stably transfected with pCDU4, harbouring the homology block E, show an overall morphology which is similar to that of wild type N4's (a cell body with neurite outgrowth). They exhibit however an increased frequency and level of lamellae formation (figure 22b). These lamellae, which contain

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F-actin microspikes are found on both the cell body and the neurite outgrowth (figure 23b). Wild type N4 cells, in contrast thereto, rarely exhibit lamellae on the neurite outgrowths.

5 3. N4 cells, stably tranfected with pCDU3, encoding for homology blocks C, D and E, show an even higher level of lamellae formation labelled with TRITC-phalloidin, the cells appear surrounded with F-actin fibres, consisting of bundles of F-actin
10 microspikes (figure 23c). The presence of these lamellae has completely modified the general appearance of the cells. They appear flatter and in 90% of the population, it is not possible to distinguish between the cell body and the wide neurite
15 as they flow gradually into one another (figure 22c). If wild-type-like thin neurite-like outgrowths are present, they are frequently numerous, branched and located all around the cell.

20 4. The overall morphology of N4 cells, stably transfected with pCDU2, encoding for homology blocks A, B, C, D, and E, resembles that of the wild type cells since, cell body and neurite outgrowth can be clearly distinguished. The pCDU2 transfected cells however show more neurite outgrowth, and these are
25 long and very branched, especially at the end of the outgrowth. When neurite outgrowths of different cells make contact, increased branching can be observed, giving the appearance of a network (figure 22d). N4 cells, transfected with pCDU2, show bundles of long
30 radial F-actin filaments (microspikes), which can be branched, especially apically. The space between the hand-shaped actin spikes is mostly filled in with actin, leading to small lamellae-like structures. Also the network-like branching between the cells
35 shows both the bundled actin structures and the

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lamellae-like fill-in features. These dense F-actin structures are sometimes seen on the cell body, which enhances the network-like appearance of the cells (figure 23d).

5 5. N4 cells, stably transfected with plasmid pTB72, encoding the full length C. elegans UNC53 protein, seem to have a more rigid structure than wild type cells, most clearly seen as spindle-like and triangle-like cells. The corners of these cells show
10 an increased level of hand-like lamellae structures. This specific phenotype is best seen when the cells are grown at low density (figure 22e, Fig. 23e).

15 Phenotypic changes in human breast carcinoma MCF-7 cells

MCF-7 cells were stably transfected with the pTB72, pCDU2, pCDU3, pCDU4 and pCB201. The population
20 of clones transfected with the LacZ-expression vector were homogeneous and similar to wild type MCF-7 cells. In contrast thereto, ~30-50% of the clones transfected with pTB72, pCDU2, pCDU3, pCDU4 and pCB201 had distinct phenotypes which were analysed as above for
25 the N4 cells:

1. Wild type and mock (pcDNA3) transfected MCF-7 cells are heteromorph. In general they are round cells or clusters of cells surrounded by lamellae. Bulges, similar to thick filopodia, can be observed
30 (figure 19a). When the cells are stained with FITC- or TRITC coupled phalloidin, F-actin actin stress fibres can be observed, often in rings surrounding the cell body (figure 20a & 21a). When cells are round up like this actin is present at the edge of the cell
35 body. Less than 10% of the cells display filopodia

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filled with radial F-actin microspikes. In time-lapse analysis the cells are highly quiescent with limited ruffling at the edge of the cell.

2. MCF-7 cells transfected with pCDU4, encoding
5 for homology block E, show two major phenotypic
differences compared to the wild type cells. These
cells are more flat and have more extended
lamellipodia leading to a pancake-like appearance.
Some clones show more filopodia than wild type (figure
10 19b). Radially organised F-actin fibres can clearly
be observed in the lamellae surrounding the cells.
These stress fibres resemble the wild-type structures,
but have a more radial than circular orientation. In
the filopodia, one can observe an increase of
15 apparently unorganised, bundles of actin patches
(figure 21b).

3. MCF-7 cells, stably transfected with pCDU3,
encoding the homology blocks C, D, and E, shows a
strikingly different and constant morphology. The
20 cells appear smaller than wild type because they are
more rounded up. All the cells have more filopodia,
surrounding the cell body (figure 19c).
Morphologically these filopodia have the same "hand-
like" appearance as those observed in N4 neuroblastoma
25 cells. Such filopodia are hardly ever observed in
mock transfected MCF-7 cells. These filopodia are
filled with F-actin fibres. Compared to wild type
cells, fine actin stress fibres are decreased (figure
21c). In time-lapse analysis single cells as well as
30 clusters of cells can be seen to ruffle much more
dynamically than single or clusters of wild type
cells. The "half-life" of a filopodia outgrowth on
the cell surface is much shorter in transfected cells
and the numbers of filopodia present at any time
35 higher.

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4. Cells transfected with pCB201 (which is structurally similar to pCDU4 but human) has a phenotype that is nearly indistinguishable from that of cells transfected with pCDU3 except that the observed phenotype and ruffling activity and filopodia outgrowth is even higher than pCDU3 (figure 18).

5. The overall morphology of the MCF-7 cells transfected with pCDU2, which encodes the homology blocks A, B, C, D and E, resembles that of the pCDU3 transfected cells. The cells are more rounded up and show more filopodia than the wild type and mock transfected cells (figure 19d). The filopodia, which are all around the cell body tend to be longer, and show a difference in actin organisation. The small filopodia have the same actin bundles as seen in the pCDU3 transfected cells. In the longer filopodia, the actin bundles are more parallel, and radial to the cell body (figure 21d).

6. MCF-7 cells transfected stably with pTB72, encoding the full length UNC53 protein, are extremely rounded up, and tend to adhere more than wild type cells. The cells grow in clusters with sausage- or tube-like shapes. The presence of large extremely thin lamellae with a surface area of more than three times the central cell body forms a second morphological feature, unique for the pTB72 transfected MCF-7 cells (figure 19e). These sheets are difficult to observe under a phase contrast microscope, but are very clear when stained with phalloidin. The lamellae protrude from one side of a cell or group of cells and are filled with thin long criss-crossing actin fibres, different from "giant" wild type MCF-7 cells (figure 21e).

These experiments lead to the following set of conclusions: (Figure 47 summarises the data of the

- 50 -

domain swapping experiments in *C. elegans* unc-53)

1. Murine and human cells transfected with the Ce-UNC-53 or hu-UNC-53/1 domains show clear effects on the nature and dynamics of their motile behaviour as demonstrated by changes in the F-actin cytoskeleton (the increase in lamellipodia, hand-like filopodia and "hair-like" microspikes on the cell surface and the associated reduction of the "rings of F-actin" stress-fibres).

2. This effect is found in two cell types of different species and tissue origin: MCF-7 cells (human breast carcinoma cells of epithelial origin) and murine N4 neuroblastoma cells. pCB201, pCDU3 and pCDU4 induce in MCF-7 cells a type of filopodium which is frequent in wild type N4 cells but rare to absent in wild type MCF-7 cells, suggesting the activation by these constructs of motile behaviour which is "normal" in N4 cells but of an unusual type in MCF-7 cells. This indicates the activation of a specific downstream process as opposed to a disruption of an existing process. It is well known that some cell types prefer to migrate with filopodia and other cell types with lamellipodia.

3. Expression of pCB201, pCDU3 and pCDU4 gives qualitatively similar F-actin remodelling and increased filopodia and lamellipodia outgrowth. pCB201 and pCDU3 are however much more active in this process than pCDU4.

4. pCB201 is a much more potent activator of filopodia outgrowth than pCDU4, which is to be expected considering the large evolutionary distance between *C. elegans* and vertebrates.

5. These experiments identify homology domain E (predicted nucleotide binding domain) of UNC-53 as the

- 51 -

"domain" that activates F-actin remodelling and filopodia/lamellipodia outgrowth. Progressive addition of the aminoterminal homology A,B,C,D lead to qualitative and quantitative modulation of the phenotype present in domain E.

6. Homology domains C and D (pCDU3) "enhance the basic activity present in homology domain E (pCDU4/pCB201).

7. Homology domains B and C (pCDU2) qualitatively modify the phenotype of domain E, leading morphologically different lamellipodia formation than pCDU3 transfected cells. It is thought that lamellipodia and filopodia formation are mediated by different signal transduction pathways requiring two related but different Ras-like G-proteins RAC for lamellipodia formation and CDC42 for filopodia formation.

8. pTB72 which includes homology domains A,B,C,D,E plus an additional 700 amino acids not yet identified isolated in the human members of the family confers a more localised filopodia outgrowth and a different morphology.

9. The expression levels of pTB72 (full length C. elegans UNC-53), pCDU3, pCDU4 and pCB201 are extremely low. The observed effect is therefore unlikely to be due to dominant negative effects (such as stoichiometric depletion of other cellular components) or structural changes in the actin cytoskeleton mediated by UNC-53 or its fragments.

The data point to a multi-domain organisation in UNC-53 whereby the aminoterminal domains exert positive (e.g. pCDU3) and negative (e.g. pCDU2) control on the activity of the domain E or are leading to novel activities or the localiation of the activity in the cell (pCDU2, pTB72). Our observation that the

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nucleotide binding domains (NTB) of distantly related members of the UNC-53 family induce similar phenotypes, suggests a general role for this domain of the UNC-53 family.

5

CELLULAR ASSAYS TO IDENTIFY PHARMACOLOGICAL
MODULATORS OF UNC-53 AND COMPONENTS OF THE UNC-53
PATHWAY

10

Mammalian and human cells transfected with plasmid constructs containing unc-53 sequence of either C. elegans or of human origin were observed to display obvious, specific and similar changes in comparison to mock or untransfected parent cells. These changes relate to the functioning of the cytoskeleton, in particular the F-actin cytoskeleton, to cell locomotion and directionally cell motility and reflect UNC-53 gene family members as capable of playing an integrator function in cell motility.

The cellular tools derived through transfection and derived functional assays with these cells not only enable characterisation of the motile phenotype typically observed after introduction of unc-53 genes, they also can be easily adapted to screen for pharmacological compounds that interfere with either (1) the expression of unc-53 gene family members, (2) the cellular functioning of unc-53 transgene(s) and of components in the unc-53 signal transduction pathway.

Two classes of pharmacological modulators are envisaged.

A first class are inhibitors of UNC-53s or the unc-53 pathway(s), which revert the described phenotypic changes induced by unc-53 transgenes or

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- 53 -

aspects thereof. Such compounds are considered relevant leads to target diseases where unwanted directional motility of cells occurs such as metastasis, angiogenesis or inflammation.

5 Secondly, pharmacological stimulators are envisaged, such as compounds which induce - in non-transfected cells - phenotypes that induce or mimic (aspects of) the described 'unc-53' phenotype. Such compounds may do so by inducing or upregulating
10 expression levels of a known unc-53 gene or by activating endogenous (yet unidentified) members of the unc-53 gene family. The target application here are wound and tissue repair, in particular diseases such as neuronal regeneration and plasticity.

15 The nature of compounds envisaged can be small (organic) molecules, bio-molecules (such as peptides, sense or antisense (oligo-)nucleotides or chemical modifications thereof. Alternatively, compounds can be thought of as a series of plasmid nucleotide
20 constructs containing gene sequences in a screen for novel unc-53-unrelated genes with a similar functional effect in the cell or genes related to the unc-53 gene family or novel members of the unc-53 gene family based on sequence similarity such as for example the
25 genes in plasmids pTB72, pcDU3, pcDU4, pcDU2, pcB201, or modifications thereof such as for example epitope tagged, deletion, complementation or mutagenised nucleotide constructs.

30 The cellular assays envisaged in the claims have been exemplified for three cell lines: the human breast carcinoma cell line MCF-7, the mouse neuronal cell line N4 and the mouse fibroblast cell line NIH-3T3. Pharmacological assays are focused on
quantification of endpoints in a high throughout
35 screening mode. Many of the computer aids for

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(semi-) automation are well known to the field and currently applied in the applicants labs. Given the subtlety of the phenotypes observed, primary focus was given to morphological assays that assess the phenotypes or aspects thereof.

The nucleotide binding domain of Hu-UNC-53/1 has transforming activity in NIH3T3 fibroblasts

10

Biochemical and genetic analysis suggest that UNC-53 functions in GRB-2 mediated signal transduction pathways controlling cell motility. The occurrence of an altered hu-UNC53/1 mRNA pattern in cancer cell lines, moved us to investigate if whether hu-UNC53/1 plays a role in the transformed state of those cells.

15

Thereto, we tested the ability of the nucleotide binding domain of hu-UNC-53/1 and Ce-UNC-53 to transform NIH/3T3 cells. Construct pCB201 (hu-UNC-53), which induces ruffling behaviour and cell motility, were transfected into NIH3T3 cells. Positive controls included Myc and H-ras. Negative controls included empty vector and Rac 1N17 and cdc42N17.

20

The cells that survived G418 selection were assayed for loss of contact inhibition (their ability to grow as foci). Positive controls included the combination of two well known oncogenes Myc and H-ras which were able to produce a high number of foci. The nucleotide binding domains of both Ce-UNC-53 and hu-UNC-53/1 are able to induce foci in this assay (Fig 24 & Table 1).

30

35

Table 1 Foci formation in NIH-3T3 cells stably transfected with pcB201

| | mock | pcB201 |
|---|------|--------|
| 5 | 22 | 138 |
| | 59 | 143 |

10

This suggests that the function of UNC-53 is not restricted to the activation of motility. UNC-53 may exert this additional function through the activation of as yet to be identified signal transduction pathways. Oncogenes frequently arise when a "controlling" domain and "activation" domain are separated through chromosomal rearrangements or integration of a part of a gene in the oncogenic virus. E.g. Erb Receptor tyrosine kinases, Ost a nucleotide exchange factor for Rac-1.

20

Hu-UNC-53/1 is localized to chromosome 1q31.1

Clone F226 (BACH-135 (014), Genome Systems, inc) was isolated from a human genomic BAC library using pCR231 as a probe and was confirmed by sequence analysis to be derived from the hu-UNC-53/1 locus. Purified DNA from clone F226 was labeled with digoxigenin dUTP by nick translation. Labeled probe was combined with sheared human DNA and hybridized to normal metaphase chromosomes derived from PHA stimulated peripheral blood lymphocytes in a solution containing 50% formamide, 10% dextran sulfate and 2X SSC. Specific hybridization signals were detected by incubating the hybridized slides in fluoresceinated antidigoxigenin antibodies followed by counterstaining with DAPI. The initial experiment resulted in

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- 56 -

specific labeling of the long arm of a group A chromosome. A second experiment was conducted in which an anonymous probe which was previously mapped to 1p34 and confirmed by cohybridization with a chromosome 1 centromere specific probe, was cohybridized with F226. The experiment resulted in the specific labeling of the long and short arms of chromosome 1. Measures of 10 specifically hybridized chromosomes 1 demonstrated that F226 is located at a position which is 52% of the distance from the heterochromatic-euchromatic boundary to the telomere of chromosome arm 1q, and that corresponds to band 1q31. At total of 80 metaphase cells were analyzed with 72 exhibiting specific labeling (Fig. 25).

Gains of DNA sequences in 1Q31 were found in more than 10% of primary bladder tumors (Genes Chromosom Cancer 12: 213-219 (1991)). A putative tumor suppressor gene located near the locus F13B on chromosome arm 1q31-q32 appears to be involved in the pathogenesis of medulloblastoma (Int. J. Cancer 67: 11-15 (1996)). Loss of heterozygosity in this region of chromosome I has been implicated in development of human hepatoblastoma. Partial trisomies of 1q31 were found in Ewing's Sarcoma cell lines isolated from patients Cancer Genet Cytogenet 12: 1-19 (1984).

HU-UNC-53/2 is localised to Chromosome 11p15.1

DNA from clone F329 from BAC for Hu-unc-53/2 was labeled with digoxigenin dUTP by nick translation and applied in the experimental settings used for FISH of Hu-unc53/1 with F226. The initial experiment with F329 resulted in the specific labeling of the mid short arm of a group C chromosome which was believed to be chromosome 11 on the base of size, morphology

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and banding pattern. A second experiment was conducted in which a biotin labeled probe specific for the centromere of chromosome 11 (D11Z1) was cohybridised with clone F329. This experiment
5 resulted in the specific labeling of the centromere in red and the mid short arm in green of chromosome 11. Measurements of 10 specifically labeled chromosomes 11 demonstrated that F329 is located at a position which is 65% of the distance from the centromere to the
10 telomere of the chromosome 11p, an area which corresponds to band 11p15.1. A total of 80 metaphase cells were analysed with 72 exhibiting specific labeling.

15 Chromosome 11p15 is a region showing loss of heterozygosity (LOH) in a variety of human malignancies, primarily breast cancer (Ali et al., Science 238, 185-188 (1987); Winqvist et al., Cancer Res. 53, 4486-4488 (1993)) but also Wilms' tumor
20 (Dowdy et al., Science 254, 293-295 (1991); Cowell et al., Br.J.Cancer 67, 1259-1261 (1993)), ovarian and testicular malignancies (Lothe et al., Genes Chromosomes Cancer 7, 96-101 (1993); Weitzel et al., Gynecol Oncol. 55, 245-252 (1994)) stomach cancer
25 (Baffa et al., Cancer Res. 56, 268-272 (1996)), lung cancer (Ludwig et al., Int.J.Cancer 49, 661-665 (1991); Fong et al., Genes Chromosomes Cancer (1994)), infantile tumors of adrenal and liver (Byrne et al., Genes Chromosomes Cancer 8, 104-111 (1993)). Since
30 LOH is believed to indicate inactivation of a tumor suppressor gene at the location where LOH occurs, the frequent LOH found at 11p15 in multiple human cancers suggests the presence of either a cluster of tumor suppressor genes or a single tumor suppressor in this
35 region (Seizinger et al., Cytogenet. Cell genet. 58,

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10080-10096 (1991)). Chromosome transfer studies have shown that chromosome 11 can suppress tumorigenicity of both human breast cancer (Negrini et al., Cancer Res.55, 3003-3007 (1995)) and Wilms' tumor cells (Dowdy et al., Science 254, 293-295 (1991)) and a gene (named HTS1 or ST5) that may be responsible for suppressing tumorigenicity in HeLa cells has been mapped to 11p15 (Lichy et al., Cell Growth Diff. 3, 541--548 (1992)). Abnormalities at 11p15 have also been identified in a variety of other cancers, including lung cancer (parental origin of 11p15 deletion) (Kondo et al., Oncogene 9, 3063-3065 (1994)), bladder cancer (Presti et al., Cancer Res. 51, 5405-5409 (1991)), myeloid leukemia (translocation) (Nakamura et al., Nat. Genet. 12, 154-158 (1996)), malignant astrocytomas and other primitive neuroectodermal tumors (deletions) (Fults et al., Genomics 14, 799-801 (1992)), rhabdomyosarcoma (Scrabble et al., Nature 329, 645-647 (1987)) and hepatocellular carcinoma (Fujimori et al., Cancer Res. 51, 89-93 (1991); Wang et al., Cell Genet. 48, 72-78 (1988)). Recently a gene, TSG101, was cloned that is mutated in human breast cancer and deleted in uncultured primary human breast carcinomas (Li et al., Cell 88, 143-154 (1997)).

DIAGNOSTIC ASSAY USING THE DNA SEQUENCE OF HUMAN UNC-53S

The differential expression of human unc-53 transcripts in Northern blots of normal tissues versus transformed cell lines and the chromosomal locus of hu-unc-53/1 at 1q31 being a locus linked to three diseases, suggests the potential implication of hu-unc-53 genes in oncogenesis. By using the complete

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DNA sequence of hu-unc-53/1 or /2 or fragments thereof in FISH, the potential involvement of these genes can be diagnosed in patients as exemplified in figure 26. Alike, the use of these hu-unc-53 sequences in
 5 diagnostic PCR assays can be used to determine overexpression of hu-unc-53s or fragments thereof.

Assay for microscopic phenotypic UNC-53
 transfected MCF-7 cells

10

Mock and unc-53 transfected MCF-7 cells were seeded at low density in culture plates and allowed to adhere to the vessel. Light microscopic inspection at different time points either on live cells or after
 15 chemical fixation with Karnovsky's fixative revealed that in pcB201, MCF-7 transfected cultures a rounded shaped cell body with at their boundaries many filopodia. In contrast, mock or untransfected clones had a predominant 'flat' phenotype - with little or no
 20 filopodia. Quantitative measurements confirmed the statistical significance of this shift in phenotype (table 2 below).

TABLE 2

25

Quantification of phenotypic changes in unc-53 transfected MCF-7 cells (*)

| Transfection: | clone | no feet (**) | with feet (**) | fraction with feet |
|---------------|-------|--------------|----------------|--------------------|
| mock | e | 34 | 8 | 0.19 |
| | | 37 | 0 | 0 |
| pcB201 | 2 | 17 | 92 | 0.84 |
| | | 37 | 83 | 0.69 |
| | 16 | 27 | 62 | 0.70 |
| | | 20 | 71 | 0.78 |
| | | 13 | 85 | 0.87 |

30

(*) Clones were passaged thrice, frozen and stored.

- 60 -

Thawed cells were trypsinised at confluency, monodispersed, seeded in flasks and allowed to attach to substrate overnight to 48 hours. Cultures were fixed with Karnovsky fixative and inspected using
5 phase contrast microscopy. In parallel experiments, resistance to genitacin was confirmed.

(**) values are expressed as cells per microscopic view.

10 Assay for ruffling and motile behaviour using automated time lapse

The dynamic changes in cells are well known in the field. Animations of e.g. actin ruffles in
15 astrocytoma cells or od actin based cell motility in e.g. fibroblasts can be accessed
(<http://www.stc.cmu.edu/CLMIBhp/Imggallpg/Moviespg/actinruffle.mov>) or
(<http://util.ucsf.edu/mitchi/Movies/migration.html>) on
20 the world wide web. The dynamic changes as a result of transfection with unc-53 can best be appreciated in time lapse video sequences. At high magnification, the 'filopodia' display arrays of microspikes with highly dynamic behaviour. A rough visual estimate
25 suggests these phenomena to be at least 10-fold increased in pcB201 transfected cells relative to the mock-transfected MCF-7 cells. Animations of these clones in NIH-Image can be requested from author or applicant.

30 Time lapse video imaging probably is the most informative way to appreciate the unc-53-induced phenotype in MCF-7 and is amenable to high throughput screening in a pharmacological context. Time lapses compressing 5 minutes real time supply sufficient
35 information to quantitate the intensity of the motile

- 61 -

behaviour of pcB201 transfected MCF-7 cells in e.g. 12 well plates. In addition, algorithms have been described in the field which can automatically compute the 'motile area' of cells by comparing cells in two
5 images appropriately spaced in time (van laerebeke et al., 1992, cytometry, 13, 1-8).

Assay for visualising unc-53-induced F-actin
recruitment in MCF-7 cells

10

Cultures were chemically fixed, detergent extracted and fluorescently stained for F-actin (filamentous-actin) using fluorescently labeled phalloidin (Wieland et al., 1985, Int. J. Peptide &
15 protein Res, 21, 3-10) which display in a more specific way the dramatic phenotypic changes to transfection with unc-53 transgenes. By using image capturing and analysis of the F-actin patterns, image analysis algorithms well known in the field can assess
20 in an automated way, the f-actin filament positions, texture and distribution relative to the nuclear position or gravity point of the cells. Such algorithms are capable of discriminating phenotypic changes and thus also effects of pharmacological
25 inhibitors of transgene-induced phenotypes as well as compound induced unc-53 like phenotypes in mock or untransfected cells.

Phagokinesis assay for unc-53-induced
30 directionality and quantity of motility

The methods are described in the experimental section. Two cell populations with different motile behaviour in phagokinesis assays were observed. In
35 table 3 below the fraction of mock and UNC-53

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transfected MCF-7 cells that produced linear tracks in the phagokinesis assay are shown. In the mock transfected MCF-7 cells, 61% of the cells produce a round track (long and short axis less than 2-fold different) and 39% cells produced 'linear' tracks (long and short axis more than 2-fold different). pcB201 transfected MCF-7 cells produced an increase of the fraction of cells displaying 'linear' tracks to 50%. An increase in the fraction linear tracks was made for MCF-7 cells transfected with full sequence Ce-unc-53.

In addition, a significant increase of 50% in the median area of tracks of a culture vessel was observed in the pcB201 transfected MCF-7 cells versus mock transfected MCF-7 cells (Table 2). These observations suggest that pcB201 as well as pTB72 transfection into MCF-7 cells is capable of increasing *in situ* locomotion in Ce-UNC-53 MCF-7, e.g. by increasing spreading, ruffling, or other forms of non-directional motility in the 'round' population. In addition the Ce-UNC-53 transgene in MCF-7 cells drives a fraction of the MCF-7 cells from non-directional motility (round tracks) into directional migration (linear tracks). Clone 2 thus provides a tool to analyse inhibitory or stimulating effects of pharmacological compounds on directionality or quantity of cell motility in relation to UNC-53.

30

35

Table 3. Analysis of motility in phagokinesis assays

Track morphology: fraction linear tracks

| | | | | | |
|------------|---------|------------------|-------|--------|------|
| 5 | plasmid | clone | round | linear | l/r |
| | Mock | z4 | 18 | 13 | 0.42 |
| 10 | | | 17 | 11 | 0.39 |
| | | | 22 | 12 | 0.35 |
| | pCB201 | Clone 2 | 16 | 9 | 0.36 |
| | | | 13 | 13 | 0.5 |
| 15 | | | 7 | 8 | 0.53 |
| | | | 9 | 9 | 0.5 |
| Track Size | | | | | |
| 20 | Clone | average \pm SD | min | max | (N) |
| | Z4 | 1626 \pm 188 | 1444 | 2011 | (8) |
| | Clone 2 | 2326 \pm 283 | 1989 | 2816 | (8) |

25 Assays for the localisation of unc-53 in the cell
to microtubules or microtubule (+) plus ends

UNC-53s have been shown to reside on microtubules and preferentially on the microtubule (+)-ends of cells. This localisation represents an important feature of the UNC-53 family of proteins, which is rarely observed in other proteins. Absence of microtubule (+)-end binding in the protein APC following mutation has been implied in the role of APC in colon cancer (Smith et al., 1994, Cancer Res., 54, 3672). In analogy, it can be postulated that the proper functioning of UNC-53 also may depend on its specific localisation in the cell.

40 The methods used in the examples which prove the co-localisation with microtubules form a base for a series of assays for compounds which specifically

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affect microtubule (+)-end binding of UNC-53s. To the skilled eye, the typical localisation of an UNC-53 protein on microtubules can be readily recognised and thus is sufficient for the interpretation of whether the treatment with a compound has affected the localisation of this UNC-53 (or a fragment thereof). Moreover, by combining the described methods (co-localisation) - well known to one skilled in the field and exemplified by the methods in the "experimental procedures" section - one can unequivocally confirm a compounds ability of abrogating (or promoting) microtubule and microtubule (+)-end binding.

Such an assay comprises contacting a cell culture of a cell line expressing an UNC-53 with a compound in the culture conditions proper for the said cell line, followed by an incubation and finally observation of the UNC-53 (or fragment) in situ by e.g. fluorescence microscopy (for GFP-chimeras) or by fixing the cell culture and performing an immunocytochemical staining for the UNC-53 (or fragment). For the co-localisation, methods such as immunocytochemistry for the microtubules of a cell or cell line combined with either immunocytochemistry for Ce-UNC-53 or Hu-UNC-53s or fluorescent detection GFP-UNC-53 chimeras are performed consecutively.

C.elegans-UNC-53 preferentially binds microtubule plus-ends or GTP-tubulin

Biochemical characterisation of UNC-53 has shown that UNC-53 binds the SH3 binding domains of SEM-5/GRB-2 and binds F-actin *in vitro*. GRB2 has been localised to the cortex of the cell and reported to be involved in the control of cell motility. To determine the *in vivo* subcellular localisation of Ce-

- 65 -

UNC-53, we transiently transfected COS, HepG2 and MCF7 cells with pTB72, an expression construct containing the full length Ce-unc-53 cDNA. This construct was previously shown to activate cell motility in N4 neuroblastoma and MCF7 cells. This construct gives high transient expression in COS cells, high to medium levels of expression in MCF7 cells and medium to low levels of expression in HepG2 cells. To visualise UNC-53, tubulin and F-actin, transfected cells were stained with various combinations of the anti-Ce-UNC-53 mab 16-48-2, rabbit anti-UNC-53 polyclonal, anti-tubulin mab YL1/2 and fluorescently labelled phalloidin.

At high levels of expression UNC-53 co-localises with the entire microtubule cytoskeleton, but at lower expression levels UNC-53 signal is restricted to the terminal regions of the microtubules at the plus ends. Very low levels of the expression yield a dot-like pattern in the vicinity of the cortex of the cell.

To map the MTB plus end domain of Ce-UNC53, we made two constructs pcDU2 (figure 17) and pcDU3 (figure 15) in which the aminotermus of Ce-UNC-53 is deleted. Proteins corresponding to these constructs are thought to be made *in vivo* from different unc-53 promoters. Transient transfections followed by immunolocalisation showed these proteins to be cytoplasmic. In stable transfections in N4 neuroblastoma cells and MCF7 cells they were shown to be no longer toxic to cells but cause highly increased activation of filopodia formation. We thus uncoupled (1) toxicity of Ce-UNC-53 from activation of motility and (2) microtubule binding from the activation of motility.

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Analysis of the microtubule association of the
C.elegans and Human 1 UNC53

To isolate the microtubule association domain of
5 the C.elegans UNC53, N-terminal GFP fusions were made.
C-terminal deletions on the fusion product revealed
that the microtubule association was localised in the
N-terminal half of the protein. A GFP fusion was also
constructed with the Human1-UNC-53, to analyse the
10 microtubule association properties of this protein.
The association with microtubules was confirmed. A
mouse anti sera was used to show the presence of
native Unc-53 on microtubule plus ends of melanoma
line G361. The epitope recognition of the antibody
15 was confirmed by immunohistology experiments with
mammalian cells, transiently expressed with pLM4,
expression the GFP-hu1-UNC53 fusion protein.

Results

20 1. When transiently transfecting pTB72 in
several cell lines C.elegans UNC-53 associates with
microtubules and preferentially the plus-ends of the
tubuline fibres. Transfection of plasmids pCDU3 and
25 pCDU2 in N4 and MCF7 cell lines did not result in the
observation of microtubule co-localisation. pCDU4
resulted in no staining using mab 16-48 antibody (LMBP
Accession No. 1383CB) concluding that the epitope for
this antibody is localised outside the fragment
30 expressed by pCDU4.

It is possible that the microtubule associated
domain is situated in the N-terminus of the protein.
For this reason, we constructed an N-terminal GFP
fusion with the full length C.elegans UNC-53 sequence,
35 and various C-terminal deletion derivatives. These

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fragments encode the N-terminal part of UNC-53 from 139 to 760 aa.

Furthermore, to analyse if the cloned fragment of hui-unc53 also could be associated with microtubules, a plasmid encoding a GFP fusion with the hui-Unc53 protein was constructed, and introduced into mammalian cells. A derivative of this construct was also constructed.

2.

a) Transient expression of C.elegans Unc-53 GFP fusion in N4 neuroblastoma lines

N4 cells were transiently transfected with pEGFP72, encoding a fusion protein of GFP and full length C.elegans unc-53 sequence. On an inverted microscope, the fluorescence of the GFP molecule could be followed in living cells. Cells which expressed low to medium levels of the fusion molecule showed a normal morphology after 18h to 30h. In these cells the co-localisation of the GFP fusion protein with the microtubules could clearly be demonstrated (figure 38a). In cells which demonstrated a low but still distinct GFP fluorescence, specific microtubule plus-end association could be observed (figure 38b). Cells expressing high levels of the GFP fusion protein tend to round up, in such a way that the microfilaments are difficult to visualise. After 48h, almost no GFP expressing cells can be found. It has previously been observed in transient expression of Unc-53, using plasmid pTB72, that the protein is toxic for the cells. The transient transfection experiments with the pEGFP72 plasmid gives the same observation, indicating that at least two features of the Unc53 protein are conserved in the GFP fusion protein, being

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the microtubule association and the toxicity of the protein.

The transfected cells were fixed with paraformaldehyde, and the tubuline was stained using antibody YL1/2 and antimouse-CY3 (Jackson Labs). Although a significant loss of GFP fluorescence was observed, one could clearly demonstrate that the filaments observed with the GFP fluorescence co-localise with the microtubules staining (figure 39).

Putative Assay

Mammalian cells, in this case N4, were transfected with a lipofecting agent (lipofectAMINE) while in suspension, not being attached to a surface. After transfecting those cells with pEGFP72, the transfected cell suspension could be diluted in 24- and/or 96-well plates, enabling them to attach to the surface. Each well may contain a different compound of the collection to screen. After 24h, plates could be automatically screened for fluorescence levels. Wells containing a compound that abolish the toxicity of the GFP-C.elegans UNC-53 fusion protein will give high levels of fluorescence. Compounds having no effect on the fusion product will give no or only low levels of fluorescence.

b) Transient expression of the truncated GFP-C.elegans UNC-53 fusion proteins.

To assay if the microtubuline association did occur in the N-terminal part of the C.elegans Unc-53 protein, various C-terminal deletions were constructed.

Transfection of pEGFPsma and pEGFPec1 coding

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for 760 AA and 670 of the N-terminal part of C.elegans UNC-53 in fusion with GFP, resulted in microtubuline association, as could be visualised in living cells. The association with the microtubules is less abundant than observed when expressing the full length C.elegans UNC-53 protein, but fibres could clearly be observed (figures 40a and 41a). More background fluorescence is seen. This could be due to a lesser association to the microtubules or to a instability of the fusion protein. The association with microtubules could not be observed after fixing the cells with paraformaldehyde nor with methanol fixation, giving an extra indication for the weak association with the microtubule network of these proteins or potential instability of the fusion protein. At low expression levels the association of the GFP fusion protein with the centrosomes could clearly be detected (Figures 40b and 41b). Centrosomes are the location in the cell with the highest microtubule concentration.

No plus-end associations could be observed with the deletion constructs, even when cells where expressing low levels of the GFP fusion proteins. In the case of very low expressions, the centrosomes could clearly be detected.

When transfecting N4 cells with pEGFPsac or pEFPXba, coding for 139 aa and 256 aa of the N-terminal part of C.elegans UNC-53 in fusion with GFP, no microtubule association could be observed. This indicates that at least 670 aa of the N-terminus of the C.elegans UNC-53 is needed to have microtubule association (figures 42a and 42b).

c) Transient expression of the GFP-hu-UNC-53/1

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fusion proteins and a deletion derivative.

Plasmid pLM4 was transiently transfected into N4 neuroblastoma cells, and GFP fluorescence was observed in living cells. GFP fluorescence of the available sequence of hul-UNC-53 in fusion with GFP was localised at the microtubule level. Moreover, at lower expression levels, both the centrosomes, and specific plus-end association could be observed. As has been observed with the C.elegans UNC-53 derivatives in fusion with GFP, expressed by the plasmids pEGFPsma and pEGFPecl, the GFP association seems to be less tight as was observed by the full length C.elegans UNC-53 fragment in fusion with GFP. The observed instability of the fusion protein can be due to a lesser association to microtubules, or to a degradation of the fusion protein (figure 43).

d) Immunofluorescence on melanoma line G361, and on neuroblastoma line N4 transiently transfected with pLM4.

Introduction

Northern experiments show that the melanoma cancer line G361 expressed abundantly both the Human1 and Human2 homologue of C.elegans UNC-53. To test if the proteins could be localised in this cell line, a collection of mouse sera was tested on this cell line. To verify if the observation was due to a hu-UNC-53 recognition, and not to an artifact, a positive sera was applied to N4 cells transiently transfected with pLM4, expressing the GFP-hul-Unc fusion.

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result

a serum, designated 28.1 from a mouse previously injected with peptide (DNRTLPPKKGLYRY) a conserved sequence of the UNC-53 family was used for a immunolocalisation experiment on G361 cells fixed with paraformaldehyde. Antimouse-cy3 was applied as second antibody. Association with microtubule plus-end could clearly be observed. Moreover, in cells showing directional movement, observed as growth cones extensions, abundant staining can be seen in the tip of the growth cone (figure 45). To test whether the recognition of the microtubule associated protein was identical to the Hui-UNC-53 protein, N4 cells were transiently transfected with plasmid pLM4 and consequently fixed with paraformaldehyde and stained with serum 28.1. Only cells that were transfected showed staining with 28.1, indicating that the antibody of 28.1 recognised the Hui-UNC-53-GFP fusion protein (figure 46). This confirms that the staining of the microtubule plus-ends in the growth cones of G361 by serum 28.1 is due to a recognition of at least the Human1 and/or the Human2 homologue. It is concluded that the overexpression of the human homologue of C.elegans UNC-53 in the melanoma cancerline G361 is located on the microtubule plus-ends.

Conclusions

30

- a) - GFP-C.elegans UNC-53 fusion protein expressed by pEGFP72 shows Unc53 activity
- b) - GFP-C.elegans UNC-53 fusion protein expressed by pEGFP72 shows microtubule association
- c) - GFP-C.elegans UNC-53 fusion protein

35

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expressed by pEGFP72 shows microtubule plus-end association

5 c) - GFP-C.elegans UNC-53-(deletion variant) fusion proteins expressed by plasmids pEGFPsma and pEGFPec1 show microtubule association.

d) - GFP-C.elegans-UNC-53-(deletion variant) fusion proteins expressed by plasmids pEGFPsma and pEGFPec1 no not show microtubule plus-end association

10 e) - GFP-C.elegans UNC-53-(deletion variant) fusion proteins expressed by plasmids pEGFPxba and pEGFPsac no not show microtubule associations.

f) - GFP-hu1-UNC-53 fusion protein expressed by plasmid pLM4 shows microtubule association.

15 g) - GFP-hu1-UNC-53 fusion protein expressed by plasmid pLM4 shows microtubule plus end association.

i) - serum 28.1 recognises the Hu1-UNC-53-GFP fusion protein as expressed by plasmid pLM4 in transiently transfected Neuroblastoma cells N4.

20 j) - the expressed human homologue of C.elegans.- UNC-53 in melanoma line (being at least hu1-Unc-53) is associated with the microtubule plus-ends.

EXPERIMENTAL PROCEDURES

25 Materials

The oligonucleotides used in the PCR-RACE experiments were synthesised by Eurogentee (Belgium). Radioactive compounds were obtained from Amersham.

30 The pCDNA3.1 eukaryotic expression vectors, human 1GT10 cDNA libraries, marathon-RACE cDNAS, human, Northern blots and the T7-tag monoclonal antibody were purchased from Invitrogen. N4, MCF7 and NIH 3T3 cells were retrieved from the Janssen Research cell bank.

35

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PCR-RACE conditions

1. A quick screen human cDNA library panel was used to amplify EST clone gb..R41071. The primers used
5 were ESTfw 5'-AATGGCTTCCTGGTTACCTGAG-3' and ESTrv 5'-CAAGTCAGCACCCCGAAGCAGCTCT-3'. Human genomic DNA was used also as template (100ng/reaction). The amplification conditions were as follows: 1 min at 94°C, 30 sec at 55°C, 30 sec at 72°C, then 35 more
10 times and a final extension of 20 min at 72°C. This PCT fragment was cloned in vector pCR2.1. The resulting plasmid was designed pCR231.

A human heart clone was also produced by RACE-PCR from a human heart Marathon cDNA using the following
15 conditions; 1 min at 94°C, 30 sec at 70°C, 3 min 30 sec at 72°C, then 35 more times and a final extension of 20 min at 72°C. KlenTaq DNA Polymerase was purchased from Invitrogen.

For the mouse homologue, total RNA was obtained
20 from N4 murine cells as described. A first strand cDNA was synthesized from 2 µgr of RNA using Ready To-Go cDNA kit (Pharmacia). The primers used were M-ESTfw 5'CCTCTGTGGGCACCGAGGTCACC--3'. The amplification conditions were as follows: 1 min at 94°C, 30 sec at
25 58°C, 30 sec at 72°C, then 35 more times and a final extension of 20 min at 72°C. All the amplifications product were subcloned in pCRII-1 and several independent clones were analyzed by sequence.

30 2. Screening of Human Heart/Colorectal Adenocarcinoma cDNA library

A human heart cDNA library and a human colorectal adenocarcinoma cDNA library were screened using
35 pCR231bp as probe by the standard plaque hybridization

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method. The screening produced several positive clones in each library called respectively λ HH3, λ HH4, λ HH15, λ CAD14 and λ CAD27. The positive phages were purified by two additional rounds of plaque screening and were then amplified.

3. 5' extension using PCR

Three primers with homology to the 5' end of clone λ HH3b were made:

HU53rv1 (5'-cct-ggg-act-gaa-gct-ggt-acc-tga-gcc-3'), HU53rv2 (5'-ttg-gga-aga-gtg-ttc-cga-tcc-cgc-tg-3') and HU53rv3 (5'gtt-gcc-cag-ctc-tgg-ggc-ttc-cac-tcc-3') and used together with λ gt10rv primer (5'-gag-gtg-gct-tat-gag-tat-ttc-ttc-cag-ggt-a-3') in three nested PCR reactions on a cDNA amplified library from Human Heart (Clontech). The reaction mixes contained 25pmol of each primer, 1 mM of each dNTP, 1 μ l KlenTaq Polymerase Mix (50x) and 0.1 ng DNA. The cycling parameters for the first PCR were: 3 min at 94°C, 35 cycles of 1 min at 94°C, 1 min at 51°C and 3 min at 72°C and a final extension of 10 min at 72°C, using HU53rv1 and λ gt10rv as primers. 0.4 μ l of this primary PCR product was amplified using HU53rv2 and λ gt10rv as nested primers with the following parameters: 3 min at 94°C, 38 cycles of 1 min at 94°C, 1 min at 52°C and 3 min 30 sec at 72°C and a final extension of 10 min at 72°C. The second nested PCR reaction was performed on 0.4 μ l of a 1/50 diluted purified 2.4 kb fragment using HU53rv3 and λ gt10rv as primers: 3 min at 94°C, 35 cycles of 1 min at 94°C, 1 min at 56°C and 3 min 30 sec at 72°C and a final extension of 10 min at 72°C. A 774 kb amplification product was subcloned in pCR2.1, resulting in plasmid pCB210-14. The clone fragment was analyzed by sequencing. This fragment

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extends 699 bp in 5' direction (see fig 9).

4. 5' extension using PCR

5 Primer HU53rv4 (5'-ccc-tgc-ttg-gtg-ctg-agg-aga-
ctg-g-3') was designed on the 5' end of clone pCB210-
14 and was used together with λ gt10rv to amplify a
fragment of the Human Heart cDNA library with the
following parameters: 3 min at 94°C, 35 cycles of 1
10 min at 94°C, 1 min at 60°C and 3 min 30 sec at 72°C
and a final extension of 10 min at 72°C. A 887 bp
fragment was subcloned in pCR2.1, resulting in plasmid
pCB212. The clone fragment was analyzed by
sequencing. This fragment extends a further 767 bp in
15 5' direction (see fig 9).

5. Human Heart Library screening using the 0.8 kb insert of pCB212 as probe

20 The EcoRI digested and purified clone pCB212 was
used as probe to screen the Human Heart cDNA library
(Clontech) using standard plaque hybridization method.
The positive phages were purified by two additional
rounds of plaque screening. The insert of the λ DNA
25 (produced using Qiagen Lambda Kit) was analyzed by
sequencing. This pHH14-3 resulted in a 2663 bp
fragment overlapping pCB212, pCB210-14 and the 3' end
(434 bp) of λ HH3b and in a 761 bp 5' extension (see
fig 9).

30

3' and 5' extension of HU-Unc53/2 from EST46037

WashU-Merck EST 46037

35 Transformed cells carrying the EST 46037 sequence
were ordered from Research Genetics. Plasmid DNA was

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isolated using standard protocols (Qiagen plasmid DNA isolation kit), the sequence of the insert was determined.

5 3' extension of EST 46037 by RACE

Marathon-Ready cDNAs (Clontech) are premade "libraries" of adaptor-ligated double-stranded cDNA ready for use as templates in RACE experiments.

10 Five ml Marathon-Ready cDNA was used as template in a regular 50ml RACE. The RACE mixture contained 1x KlenTaq PCR buffer, 0.2 mM of each dNTP, 1x advantage KlenTaq polymerase mix (Clontech), 0.15 mM AP1 adaptor primer and 0.15 mM RACE gene specific primer. The
15 amplification conditions were as follows :

94°C for 1 min, 5 cycles of 94°C for 30 s and 72°C for 4 min, 5 cycles of 94°C for 30s and 70°C for 4 min, 25 cycles of 94°C for 30 s and 68°C for 4 min.

20 One-hundred-fold diluted RACE product was used as a template in a nested PCR with AP2 adaptor and gene specific nested PCR primers. Specific nested PCR fragments were cloned into pCR Γ 2.1 (TA cloning kit, Invitrogen) and the sequences of the inserts were
25 determined.

gene specific primer (EST46037-F1)

5'AGTGAGAACAATGCTGTGGACATGC nested gene specific
primer (ES46037-F2) 5'CTGCTCAACTGCAAGTACCACAAATGC
Marathon cDNA library : human placenta

30

WashU-Merck EST 923793

Transformed cells carrying the EST 923793
sequence were ordered from Research Genetics. Plasmid
35 DNA was isolated using standard protocols (Qiagen

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plasmid DNA isolation kit), the sequence of the insert was determined.

5 RACE fragments 1.4 and 3.7, 5' extension of
 EST46037

 Method as described previously. Gene specific
 primer (EST46037-R1) 5'ACTGCCTTGAGACTCTGACTTCAGC
 nested gene specific primer (ES46037-R2)
10 5'TGGGCAGAACTGAGAGCTTCTAAGC Marathon cDNA library :
 human placenta

RACE fragments B2.1, D2.1, H2.1; 5' extension

15 Method as described previously:gene specific
 primer (97010709) 5'ATTCTTTTGCATCTTCTTGCGTGCG
 nested gene specific primer (97010708)
 5'ACCTGAGTCCTTTCTTAGGCAAAGTGTTCC Marathon cDNA library
 : human placenta (fragment B2.1)

20 human HeLa S3 (fragment D2.1) human colorectal
 adenocarcinoma SW480 (fragment H2.1)

PCR fragments E2.3, C2.3

25 EST 485068 is similar to but not identical with
 the 5'end of HU-Unc53/1. A primer pair consisting of
 one 3' EST 485068 primer and one 5' HU-Unc53/2 primer
 were used to PCR amplify those fragments. 1gt10 human
30 placenta Quick screen library (fragment C2.3) or
 Marathon cDNA from human HeLa S3 (fragment E2.3) were
 used as templates in a PCR. A 50 ml reaction mix
 contained 1xPCR II buffer (Perkin-Elmer), 1.5 mM
 MgCl₂, 0.2 mM of each dNTP, 0.15 mM forward and
35 reverse primer, 2.5 U AmpliTaq Gold (Perkin-Elmer)

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and 1 ml template. The cycling parameters were 5 minutes at 95°C, 35 cycles of 45 seconds at 94°C, 45 seconds at 65°C and 2 minutes at 72°C. The PCR products were sliced out from an agarose gel and purified using a gel extraction kit (Qiagen), one ml hereof was used in a second round PCR using the same conditions as above. The PCR products were purified (Qiagen PCR purification kit) and direct sequenced.

primers :

(97010709) 5'ATTCTTTTGCATCTTCTTGCGTGCG

(97012802) 5' CGCTCCCCATCAGATGCAGGCCGG

PCR fragment E1.3-3

EST 01222 is homologous but not identical with the 5' end of HU-Unc53/1. A primer pair consisting of one 3' EST 01222 primer and one 5' HU-Unc53/2 primer were used to PCR amplify this fragments. Marathon cDNA from human HeLa S3 was used as template in a PCR. A 50 µl reaction mix contained 1xPCR II buffer (Perkin-Elmer), 1.5 mM MgCl₂, 0.02 mM of each dNTP, 0.15 mM forward and reverse primer, 2.5 U AmpliTaq Gold (Perkin-Elmer) and 1 ml template. The cycling parameters were 5 minutes at 95°C, 35 cycles of 45 seconds at 94°C, 45 seconds at 65°C and 2 minutes at 72°C. The PCR products were sliced out from an agarose gel and purified using a gel extraction kit (Qiagen), one ml hereof was used in a second round PCR using the same conditions as above. The PCR products were analysed on an agarose gel, the fragment of interest was sliced out, purified (Qiagen PCR purification kit) and cloned into pCRIT2.1. The sequence of the insert was determined.

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RACE fragments A2.2-2, B2.1-4, D2.1-5; 5'
extension

5 Method as described previously.
gene specific primer (97041701)

5'TATGCTACGGCCACTCATCTCCGTGG
nested gene specific primer (97041702)

10

5'TGTAACCTGAGTTCCCCTTAAACTGG

Marathon cDNA library :

human placenta (fragment A2.1-2)

human HeLa S3 (fragment B2.1-4)

15 human colorectal adenocarcinoma SW480 (fragment
D2.1-5)

Translation-initiation splice variants, fragments
D4.1-1, J4.1.4, G4.1.1, F4.1.2

20

Four different translation initiation splice
variants were detected by 5'RACE.

Method as described previously.
25 gene specific primer (97080803)

5'TCGGTTGTTAGCAGTAGTTGACCCTCC
nested gene specific primer (97080804)

30 5'ACCTGAAAGTCTGGACTGCATTTTCAGC
Marathon cDNA library : human colorectal
adenocarcinoma SW480 (fragment D4.1-1) gene specific
primer (97080801)

35 5'ACAACCTGGATAATCTGGGCCAGGAGG

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nested gene specific primer (97080802)

5'TCTTGCTGGAGATCCTTGATGAGACGC

Marathon cDNA library :

5

human melanoma G361 (fragment J4.1.4)

human HeLa S3 (fragment G4.1.1)

human placenta (fragment F4.1.2)

10

DNA sequencing

PCR amplification products and cDNA clones were subcloned either into pBluescript vectors (Stratagene) or in PCR-IIa vector (Invitrogen) and sequenced either manually by the dideoxynucleotide chain termination method with modified T7 DNA polymerase (Sequenase, United States Biochemical) or automatically with an Applied Biosystems 373 DNA sequencer using the fluorescent terminator kit (Perkin Elmer).

20

RNA blots

A Human multiple tissue Northern (MTN-1, Clontech) containing in each lane 2 mg of poly A + RNA from eight different human tissues (heart, brain, placenta, lung, liver, skeletal muscle, kidney, and pancreas) and a MTN-II human multiple tissue Northern, containing in each lane 2 mg of poly A + RNA from spleen, thymus, prostate, testis, ovary, small intestine, colon and peripheral leukocyte, were hybridized according to the manufacturer's instructions and washed out in 0.1xSSC:0.2% SDS at 55°C. Also from Clontech, a poly A + RNA blot from human cancer cell lines (melanoma G361, lung carcinoma A549, colorectal adenocarcinoma SW480, Burkitt's

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lymphoma Raji Leukemia Molt 4, lymphoblastic leukemia K562, HeLa S3 and promyelocytic leukemia HL60) was tested.

5 Construction of plasmids

Plasmid pCDU2 (Figure 17) was constructed by cloning the 2.8 kb *ApaI*-*NarI* fragment from pTB72, the latter restriction site made blunt with klenow enzyme, into pCDNA3, digested with *EcoRV* and *ApaI*. pCDU2 encodes for the homology blocks A, B, C, D and E. Plasmid pCDU3 (Figure 15) was constructed by cloning the 1.9 kb *ApaI*-*NdeI* fragment from pTB72, the latter restriction site made blunt with Klenow enzyme, into pCDNA3, digested with *EcoRV* and *ApaI*, pCDU3 encodes for the homology blocks C, D and E. Plasmid pCDU4 (Figure 16) was constructed by cloning the 1.4 kb *ApaI*-*StyI* fragment from pTB72, the latter restriction site made blunt with Klenow, into pCDNA3 digested with *EcoRV* and *ApaI*. pCDU4 encodes for the homology block E.

Expression of a domain of the human UNC53 in eukaryotic cells

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1. pCB201: Equivalent construct of human 1 homologue to expression construct pCDU4 of *C. elegans* unc-53 gene cloned in a eukaryotic His-tag, Xpress Ab tag expression vector.

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A suitable Bam HI site was engineered on pHH15 open reading frame by amplification with hh15fw primer 5'AGAGCGGATCCATATGCCTCCTTGCCGTCAAGGTG-3' and M13rv primer (5'-cag-gaa-aca-gct-atg-ac-3'). The amplified fragment was then moved to pCDNA3.1.His-A-Vector

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digested with BamHI and EcoRI. This new plasmid called pCB201 (Figure 13) produces a cDNA which codes for a fusion protein consisting of a 49 amino acid aminoterminal fragment containing an His-tag and also a T7 epitope tag followed by amino acids 1255 to 1627 of the sequence of the human homologue. pCB201 was also checked by sequence and the n was used in stable transfection experiments carried out in N4, MCF7 and NIH3T3 cells.

2. pLM5: Equivalent construct of human 1 homologue to expression construct pCDU3 cloned in an eukaryotic His-tag, Xpress Ab tag expression vector.

The phage HH3b was linearized using XhoI. A BamHI and XbaaI restriction site were created on the pHH3b open reading frame using U3-Bfw (5'-cca-cac-tag-ggg-atc-cat-gca-aat-gag-g-3') and U-rv (5'-caa-aag-tct-cta-gag-gag-gcc-agt-3') as primers. This amplified fragment was then moved to pBluescript KS, digested with BamHI and XbaI. Sequencing of this plasmid, named pCB300, showed an amino acid change from a serine to an asparagine due to a change from guanine to adenine on the position 4237 of the DNA sequence. This fault was repaired by cloning a 1418 bp fragment of pLM1 (see below) (using NarI and XbaI as enzymes) into pCB300 digested with the same enzymes. The phage HH3b fragment of this plasmid, named pLM6 (fig 53), was then removed using BamHI and XbaI, to pCDNA3.1/HisA digested with the same enzymes. This new plasmid, named pLM5 (fig 52), produces a cDNA which codes for a fusion protein consisting of a 49 amino acids aminoterminal fragment harboring a His-tag and a T7 epitope tag, followed by aminoacid 1069 to 1627 of the transcript of HU-Unc53/1. Plasmid pLM5 was

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checked by sequencing and used on transient and stable transfection experiments carried out in N4 cells. The plasmid pLM1 was created using a PvuII and partial BamHI digested fragment of pHH14-3 and a BamHI and SpeI digested fragment of phage HH3b, cloned into pBluescript KS digested with SmaI and SpeI. The pLM1 contains the full transcript of HU-UNC-53/1 available at this moment (see fig 9).

3.pCB251: Equivalent construct of human 1 homologue to expression construct pCDU2 cloned in an eukaryotic His-tag, Xpress Ab tag expression vector

The phage HH3b was linearized using XhoI. A BamHI and XbaI restriction site were created on the pHH3b open reading frame using U2fw (5'-aag-gga-tga-ttc-ggt-cag-gat-cct-tc-3') and U-rv (5'-caa-aag-tct-cta-gag-gag-gcc-agt-3') as primers. The amplified fragment was then moved to pCR2.1. This plasmid was named pCB250. The pHH3b fragment was removed from pCB250 using BamHI and XbaI and cloned in pCDNA3.1/HisC digested with the same enzymes. This plasmid, named pCB251 (figure 55), was checked by sequencing. pCB251 produces a cDNA which codes for a fusion protein consisting of a 49 amino acid aminoterminal fragment harboring a His-tag and a T7 epitope tag, followed by amino acids 828 to 1627 of the partial transcript of HU-Unc53/1. pCB251 was used on transient and stable transfection experiments carried out in N4 cells (see fig 56).

4. pLM3: the partial transcript of HU-Unc531 cloned in an eukaryotic His-tag, Xpress Ab tag expression vector

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pLM1 was digested with EcoRV and XbaI. This fragment was cloned in pcDNA3.1/HisB, digested with the same enzymes. pLM3 produces a cDNA which codes for a fusion protein consisting of a 49 aminoacid aminoterminal fragment harboring a His-tag and a T7 epitope tag, followed by amino acids 1 to 1627 of the transcript of HU-Unc53/1 available at this moment. pLM3 was used on transient and stable transfection experiments carried out in N4 cells.

5. pLM4: the partial transcript of HU-Unc53/1 cloned in an eukaryotic GFP expression vector

pLM1 was digested with ClaI and XbaI. This fragment was cloned in pEGFP-cl, digested with AccI and XbaI. This plasmid was named pLM4. This plasmid produces a cDNA which codes for a fusion protein consisting of GFP, followed by aminoacid 1 to 1627 of the transcript of HU-Unc53/1. pLM4 was used on transient and stable transfection experiments carried out in N4 cells (see figs 43 and 46).

Stable transfection of MCF-7 cells:

Cells were seeded at a density of 2×10^6 cells in a 75 cm² flask using standard culture medium ((Dubecco's MEM, 450 mg/l glucose, 862 mg/l L-Alanyl-L-Glutamin, 110 mg/l Na-pyruvate; GibcoBRL) supplemented with 10% foetal calf serum (FCS; GibcoBRL), and 100 U/ml penicillin (GibcoBRL) and 100 µg/ml streptomycin). The culture was grown at 37°C in a 10% CO₂ atmosphere, to approximately 70% confluency (approximately 18 hours). The culture medium was removed and 10 ml MEM-HEPES (GibcoBRL) supplemented

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with 10% FCS was added to the cells. The culture was further incubated for four hours at 37°C in standard sterile air. DNA-CaCl₂ was meanwhile prepared by mixing 30 µg DNA in 0.1 x TE (1 mM Tris. HCl, pH 7.2, 0.1 mM EDTA, pH 8) and 0.1 ml 1.25 M CaCl₂/HEPES (1.25 M CaCl₂, 0.125 M HEPES; pH 7.05). 0.1 x TE was added to a final volume of 0.5 ml. The DNA-CaCl₂ was added drop by drop to 0.5 ml BS/HEPES (25 mM HEPES, 0.25 M NaCl, 0.01 M KCl, 1.4 mM Na₂HPO₄, 0.01 M glucose, pH 7.05) while pipeting a sterile airflow through the latter solutions. The DNA-Ca₃(PO₄)₂ precipitate was then placed at 37°C for ten minutes. The DNA-Ca₃(PO₄)₂ precipitate was vortexed and added to the cells, together with 100 µl of a 0.01 M chloroquine (Sigma) stock in H₂O. After four hours of incubation at 37°C in sterile standard air, the medium was removed, and the cells were washed with PBS (GibcoBRL). 25 ml of medium was added and the cells were placed at 37°C in a 10% CO₂ atmosphere. After 48 hours of incubation, the cells were harvested, diluted and cultivated under selection (600 µg/ml G418 (Duchefa)) for two weeks prior to clone selection. Mock transfected MCF-7 were positive for the beta-galactosidase transgene. The stability of transfection in MCF-7 was assessed by passaging cells four times in the absence of Geneticin and then re-exposing them to the selector agent. In these experiments, unc-53 or mock transfected cells proliferated, whereas untransfected MCF-7 cells proliferated at a much slower rate.

Stable transfection of N4 neuroblastoma cells

Cells were seeded at a density of 2x10⁶ cells in a 25 cm² flask using standard culture medium ((MEM Rega 3; GibcoBRL) supplemented with 10% FCS, 0.14%

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Na₂CO₃,
2 mM glutamine, 100 U/ml penicillin, and 100 µg streptomycin). The culture was grown overnight at 37°C in a 10% CO₂ atmosphere. Transfection mixture was prepared by adding 12 µg DNA in 600 µl opti-mem 1 (GibcoBRL) to 36 µl LipofectAMINE (GibcoBRL) in 600 µl opti-mem 1. This was done by adding drop by drop the first solution to the second. The mixture was placed for 30 minutes at room temperature, after which 1.8 ml of opti-mem 1 was added. In the meanwhile the cell culture was washed twice with opti-mem 1, and the 3 ml of transfection mixture was added. The culture was placed at 37°C in sterile standard air. After four hours, 3 ml of normal culture medium was added and the culture was placed at 37°C under 10% of CO₂. 18 hours later, the culture was washed with PBS, and fresh normal culture medium was added. A further 24 hours later, the cells were harvested, diluted and cultured under selection (750 µg/ml G418) for two weeks prior to clone selection.

Fixation of cells for Immunofluorescence

Medium was removed from the 9 cm² wells containing the coverslips. A 4% solution of paraformaldehyde (Sigma) in PHEM (1 g/l glucose, 0.4 g/l KCl, 8 g/l NaCl, 0.06 g/l KH₂PO₄, 0.0475 g/l Na₂HPO₄, 0.35 g/l NaHCO₃, 1.51 g/l PIPES, 0.76 g/l EGTA, 0.19 g/l MgCl₂; pH 6) was added for 30 min at room temperature. The fixative was removed, and the coverslips were washed three times 10 minutes with PHEM. The coverslips were then placed in PHEM, containing 0.5% Triton-X100 (Serva) for 30 minutes, after which the slide was washed again for three times 10 minutes with PHEM. The coverslips were then placed

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under PBS (0.14 M NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, 1.8 mM KH₂PO₄, pH 7.3) containing 0.2% Tween (Sigma) for at least one hour at 4°C

5 Immunofluorescence staining

 The coverslips were inverted on 35 µl of appropriately diluted antibody, being YL 1/2 for tubulin and/or mab 16-48-2 monoclonal or anti-UNC53
10 (gp48) polyclonal antibody for UNC53. The slides were placed at 4°C for at least 18 hours. Excess of primary antibody was then removed by washes of three times ten minutes in PBS-Tween. The slides were then treated with secondary antibody in the same way as for
15 the primary antibody. F-actin was labelled by including TRITC- or FITC coupled phalloidine to the incubation buffer. The inverted slides on the secondary antibody were left at room temperature for approximately one hour. Slides were then washed again
20 for three times ten minutes with PBS-Tween and once with PBS. The coverslips were mounted on slides with the medium described by Herzog et al. (Cell Biology: a laboratory handbook, 1994, Academic Press, 355-360). After at least two hours, slides were ready for
25 analysis.

Time lapse analysis

 Analysis of the behaviour and movement of growing
30 cell cultures was done by placing a non-confluent culture under a phase contrast microscope equipped with a temperature controlled stage (37°C). Images were recorded using a CCD camera (COHU 4912) coupled to a SCION LG3 framegrabber in a Macintosh ppc 8100
35 running NIH image version 1.60. Images were recorded

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at time intervals, varying from 15 sec to 1 min. for half an hour to two hours. Image enhancement and playback was done in NIH image.

5 Phagokinesis

A variety of cell types were shown to migrate over colloidal gold coated culture plastic or glass and displace or phagocytose the gold lawn on their way while locomoting. The track left bare is a qualitative and quantitative measure of cell motility and/or locomotion. The basic methods have been described in detail elsewhere (Albrecht-Buehler, 1977, Cell, 11: 395, Zetter, 1980; Nature, 285: 41; O'Keefe et al., 1983; J. Invest. Dermatol., 85: 130). Culture plates were gelatin and gold coated as described by Albrecht-Buehler (1977). Unc-53 and mock transfected MCF-7 were seeded in plates at low density and allowed to adhere to the plate and to locomote overnight. Cells were chemically fixed to the plate, washed and air-dried. Images of the gold lawns were captured using automated videomicroscopy; composite images of the wells were generated and single-cell phagokinesis tracks were measured using a home-made routine in SCILTM software.

C. elegans-UNC-53 preferentially binds microtubule plus ends or GTP-tubulin

1. Cloning of C.elegans cDNA in pEGFP-C1 and construction of C-terminal deletion derivatives.
 - a) Constructing a GFP-Unc53 N-terminal fusion:
A PCR experiment was performed under standard conditions, using pTB72 as template and cp17

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(ata gcc aga tct acg tca aat gta gaa ttg) and cp18
(ttt aga aac cgc ggg tgg) as primers. The resulting
0.4 kb fragment, coding for the N-terminal fragment of
C.elegans Unc 53 was cloned in vector pCR2.1 (original
5 TA cloning kit, Invitrogen), resulting in plasmid
pTA1718. The 0.4 kb fragment was isolated as a BglIII-
SacII fragment and cloned in pEGFP-C1 (Clontech)
digested with the same enzymes. The resulting plasmid
was designated pEGFPsac (Figure 29). pEGFPsac encodes
10 the N-terminal 13 aa of C.e.Unc53 in fusion with GFP.

b) Construction of a GFP-C.e. Unc53 full length
fusion:

Plasmid pTB72 (shown in Figure 1) was
digested with restriction enzymes SacII and ApaI. The
15 resulting 4.5 kb cDNA fragment, encoding for the C-
terminal fragment of C.elegans Unc53 was cloned in
plasmid pEGFPsac (Figure 29), digested with the same
enzymes, resulting in plasmid pEGFP72 (Figure 30).
Plasmid pEGFP encodes GFP in fusion with the full
20 length C.e. Unc53.

c) Construction of N-terminal deletions of GFP-
C.elegans UNC-53 fusion protein, other than pEGFPsac:

pEGFP72 was digested with SmaI. The
resulting 7.0 kb fragment was religated and
25 transformed in E.coli, resulting in plasmid pEGFPsma
(Figure 31). This plasmid codes for the first 760 aa
of the Ce-UNC-53 in fusion GFP.

pEGFP72 was digested with restriction enzymes
Ecl136II and SmaI, the resulting plasmid after
30 ligation and transformation in E.coli of the 6.7 kb
fragment was designated pEGFPec1 (Figure 32). This
plasmid codes for the N-terminal 670 aa of the C.e.
Unc53 in fusion with GFP. pEGFP72 was further
digested with SmaI and XbaI. The latter site was made
35 blunt with Klenow polymerase. The resulting fragment

- 90 -

of 5.4 kb was religated and transformed in E.coli. The resulting plasmid was designated pEGFxba (Figure 33). This plasmid codes for the N-terminal 256 aa of C.elegans Unc53 in fusion with GFP.

5

2. Constructing a hul-UNC-53-GFP fusion, and a deletion derivative

The 5.4 kb hul-unc53 fragment was isolated as ClaI-XbaI fragment from pLM1 (Figure 54), and cloned in pEGFP-C1 digested with AccI and XbaI. pEGFP-C1 was isolated from E.coli GM41 (Hfa H, dam-3, thi-1, rel-1). This makes the XbaI restriction site available for restriction digest. The resulting plasmid was designated pLM4 (Figure 34).

15

3. Visualisation of GFP fluorescence in N4 cells

N4 neuroblastoma lines were seeded in Lab Tek chambered coverglass (Nalge Nunc International) and transfected using lipofectAMINE (GibcoBRL). After 18 hours, the chambered coverglasses were placed on an inverted microscope, and GFP fluorescence could be visualised.

25

4. Staining GFP fusion expressing cells with antibodies

Transfection with the GFP fusion constructed was also performed on coverglasses in a 6-well plate. After paraformaldehyde or methanol-acetone fixation, cells could be stained for actin cytoskeleton with TRITC-phalloidine, for hul-unc53 with sera 28.1 and for tubuline with YL1/2 antibody. Visualisation was then

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performed on a axioplan (Zeiss microscope).

5 Methods of Producing and Observing the Effects of
 A Chimeric unc-53 Gene

1. Definition of a promoter region in the unc-53 C.elegans gene:

10 The genomic region from the position 15621 to
 18415 in the C.elegans unc-53 gene, called promoter A,
 was cloned and fused to the cDNA of the GFP gene
 (clone pA/GFP, or pNP10)(cf. fig.51). This construct
 is injected into wild type worms (N2). Transgenic
15 line express GFP in different neurones: the two pairs
 of pioneering neurones PVP and PVQ, both BDU neurones,
 both ALN and PLN neurones, both PDE neurones, both PVM
 neurones, and 4 vulval cells. Expression begins in
 early embryogenesis, when the axons of those neurones
20 grow out.

2. Mutant Phenotype in Unc-53(n152) alleles:

25 In wild type worms (N2), the two pairs of ALN and
 PLN neurones each send an axon in a straight line
 longitudinally from the tail to the head (see
 fig.50a). In unc-53(n152) alleles, the axons are
 shorter and often branch in a dorso ventral direction
30 (see fig.50b). The neurones are visualised with the
 construct pA/GFP, injected in unc-53(n152) worms.

3. The minigene pA/unc-53 rescues the
 elongation defect of ALN and PLN neurones:

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The promoter A from the *C.elegans* unc-53 gene was fused to the cDNA of the *C.elegans* unc-53 gene (clone pA/unc53, or pNP9). This construct was injected in unc-53(n152) mutant worms, together with the pA/GFP construct described above to visualise the ALN and PLN neurones. The elongation defects of those neurones in the unc-53 mutant are almost completely restored by the expression of the unc-53 cDNA express under the promoter A (see figs. 50 and 51b).

4. Domain swap between the *C.elegans* and human unc-53 gene:

To test whether the vertebrate and worm members of the unc-53 family are functionally equivalent, we tested the ability of the human gene to rescue the mutant phenotype in the worm. We replaced the carboxyterminal predicted nucleotide binding domain (NTPase) of the worm protein with the homologous fragment of the human 1 gene.

The clone pA/unc-53 was deleted of the *C.elegans* NTPase domain, from the HpaI site, position 29800 on the genomic of unc-53, and replaced by the equivalent domain of the human-1 gene (unc-53H1) (see fig. 51). The resulting clone is named pA/unc-53H1. When this clone is injected to unc-53(n152) mutants, the transgenic worms show a significant but incomplete rescue of the defect in the elongation of the ALN and PLN neurones (see fig. 51b). The axons are longer, often elongated until the region of the vulva in a straight line, without branching dorsally anymore. This result shows that a NTPase region of the human unc-53 homologue can functionally replace the NTPase region of the *C.elegans* worm.

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The degree of rescue was analyzed quantitatively and summarized in Figure 51b:

The four strains compared are:

wt; un-53(n152); unc-53(n152),pA/unc-53;unc-

5 53(152),pA/unc-53-H1.

The various phenotypes observed are brought together in three large classes:

<<wild type>> the axon is straight, unbranched and migrates into the head;

10 <<vulva>> the axon is straight, unbranched and stops in the vulva region;

<<mutant>> the axon is short, does not reach the vulva region and has collateral branches.

The figures are indicated as a percentage. The number
15 of axons observed is indicated in the following column.

The data clearly show demonstrate conclude that the nematode/human chimera minigene pA/unc-53-H1 partly rescues the defects of the axonal migration of
20 the ALN and PLN neurones and demonstrate conservation of function of this domain between man and worm. The transgenic lines provide a functional screening assay for the motility function of at least part of the human UNC-53 gene.

25

II. Materials and methods

1 - Cloning:

30 a) pAB/GFP (pNP3 - Figure 27)

The gene of GFP has been amplified by PCR with cpn3 oligo-nucleotides

"acattaagcttcgtacgcttgagggtaccg" and Cpn5

35 "gaaaggatccgtacgataaggtattttgtgtcgg" on the plasmid pPD95.75(Figure 59) so as to be inserted at the 5'

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position in fusion into the exon 12 of the unc-53 gene at a single restriction site *Spl*I and contains its stop codon at 3' plus one polyadenilation site. The PCR amplification product is directed by *Hind*III and *Bam*HI, sites which are contained respectively in the *cpn3* and *cpn5* oligonucleotides and sub-cloned in the pBS vector (clone pNP2). The GFP is then excised from the pNP2 clone at the site *Spl*I and integrated into the X16 clone (Figure 60) originating from sub-cloning of the lambda phage S4 digested by *Xho*I. The X16 clone containing the genomic sequence of unc-53 from the position 16621 to the position 24891 cloned in the site *Xho*I of pBS.

b) pAB/unc-53 (pNP8 - Figure 35)

The promoter region AB of the X16 clone (between *Pst*I and *Spl*I) has been inserted in the clone pTB115 (Figure 58) in which the region between the sites *Pst*I and *Spl*I, containing the promoter of the gene *mec-7* and the start of the gene unc-53, has been removed.

c) pA/GFP (pNP10 - Figure 56)

The promoter region A has come from the X16 clone between the sites *Pst*I and *Nhe*I and integrated in the vector pPD95.75 containing the GFP in the sites *Pst*I and *Xba*I.

d) pA/unc-53 (pNP9 - Figure 44)

The promoter region A has come from the X16 clone between the sites *Pst*I and *Bst*XI and is integrated into the clone pTB115 in which the region between the sites *Pst*I and *Bst*XI, containing the promoter of the gene *mec-7* and the start of the gene unc-53, has been removed.

e) pA/unc-53 -H1 (pCB501 - Figure 57)

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The clone pA/unc-53 (pNP9) has been deleted from the region 3' of the gene unc-53 of the nematode between the sites HpaI and NcoI. The 3' region of the Hlunc-53 gene has been amplified by PCR with the oligonucleotides U4Afw (5'-gca-cat-cgt-taa-cgg-gga-
5 ctt-gaa-gc-3') and Urv (5'-caa-aag-tct-cta-gag-gcc-agt-3') and digested with HpaI and XbaI. After a filling stage with T4 polymerase, the ligation is effected with a complete end.

10

2-Injection

Conventional injection techniques are used (Fire A., 1986, Mello G, et al, 1991, journal Mello G. and Fire A., 1995). Young hermaphrodite adults are injected in their two syncytial gonads. The DNA used is prepared in standard manner (Qiagen) followed by precipitation with lithium chloride. After an extensive rinsing stage to eliminate all the salts, the DNA is
20 resuspended in water. The injection solution contains the different DNAs at a concentration of 100 ng/μl in an injection buffer. The plasmid pRF4 containing the dominant allele su 1006 of the gene rol-6 (Kramer J. et al, 1990, Mello C. et al, 1991) is used as a
25 transformation co-marker. The descendants of roller phenotype of the hermaphrodite injected are isolated. Approximately 10 % of these transformants will yield a stable strain, in which the different DNAs injected are associated to form a mini-chromosome which will
30 segregate as unstable extrachromosomal arrays. All the transgenic strains obtained were tested by PCR for the presence of the DNA injected, using a specific primer of the vector and a primer in the gene (results not shown).

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3. Microscopy

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The nematodes are observed under a ZEISS Axioplan microscope provided with Nomarski lenses, with 40X Neofluar, 63X Plan-Apochromat, 100X Plan-Apochromat objective lenses. For fluorescence observation the luminous source is a mercury bulb. Different ZEISS filters are used:

- for observation under GFP fluorescence, FITC filter: blue excitation line at 588 nm, emission through a 515-565 nm band-pass filter;

- for observation of the antibody labelling with a secondary antibody coupled to the TRITC: excitation through a 546 nm band-pass filter, emission through a 590 nm long-pass filter.

The image acquisition is effected by means of a CCD camera and and NIH image program using a Machintosh computer. The images are processed using the Adobe Photoshop program.

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Sequence Listing

The following sequences are referred to in the specification:

5

Sequence ID No 1 is an amino acid sequence of human homologue 1 of UNC-53 protein illustrated in Figure 9b.

10 Sequence ID No 2 is an amino acid sequence of human homologue 2 of UNC-53 protein illustrated in figure 11d.

Sequence ID No 3 is a nucleic acid sequence of the hu-1-unc-53 gene illustrated in Figure 9b.

15 Sequence ID No 4 is a nucleic acid sequence of the hu-2-unc-53 gene illustrated in Figure 11d.

Sequence ID No 5 is a nucleotide sequence of Phage Lamda Clone 3b deposited under Accession No LMBP 3595.illustrated in Figure 9.

20 Sequence ID No 6 is a nucleotide sequence of plasmid pLM1 deposited under Accession No LMBP 3762 and illustrated in fig 54.

Sequence ID No 7 is a nucleotide sequence of plasmid pLM4 deposited under Accession No 3763 and illustrated in fig 34.

25 Sequence ID No 8 is a nucleotide sequence of plasmid pEGFP72 deposited under LMBP Accession No 3764 and illustrated in fig 30.

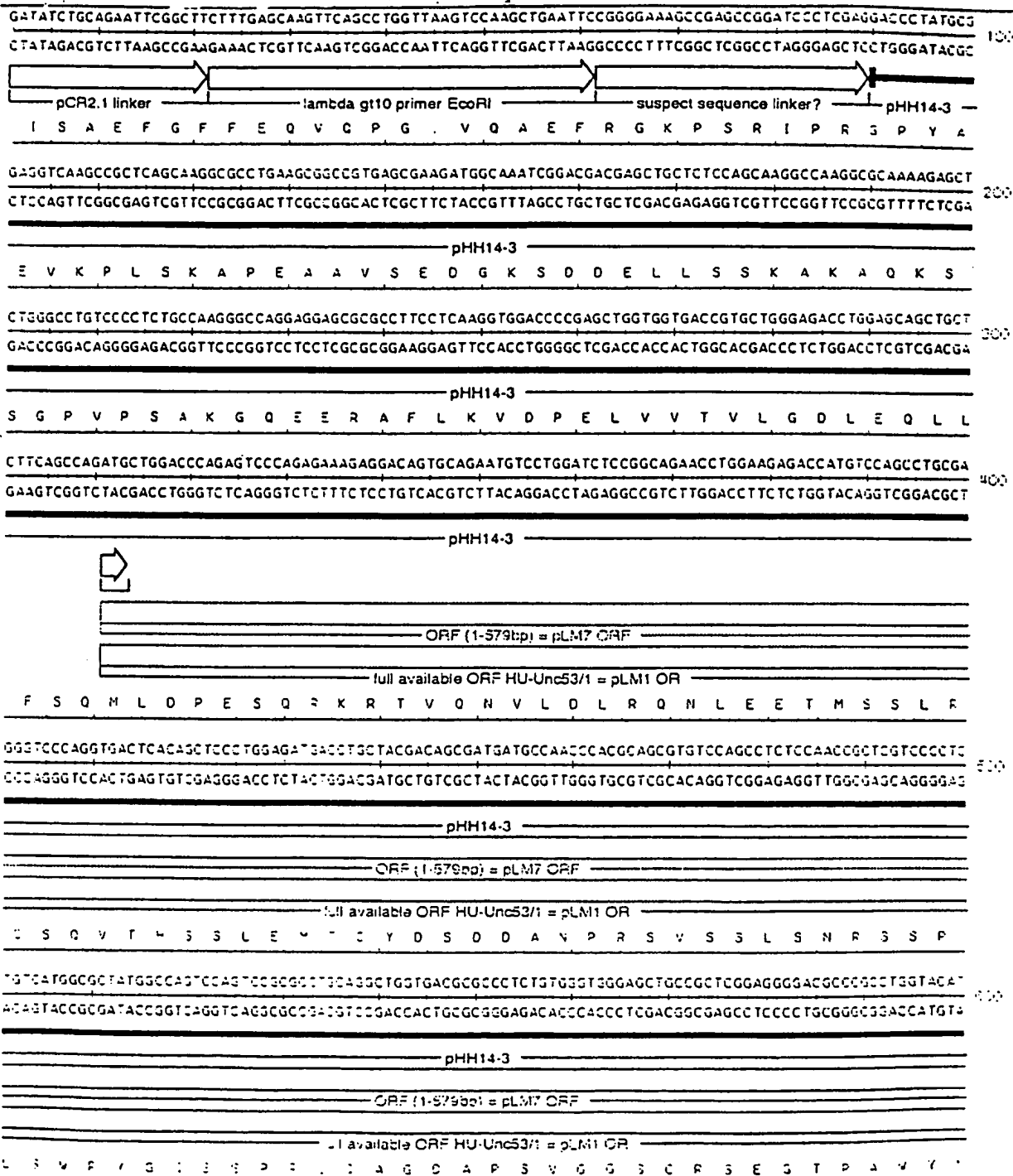
30 Sequence ID No 9 is a nucleotide sequence of plasmid pCB501 deposited under Accession No 3765 and illustrated in fig 57.

Sequence ID No 10 is a nucleotide sequence of plasmid pCB201 deposited under Accession No. LMBP 3594.

SEQ ID NO. 1

FIND ACID SEQUENCE OF
HU-UNC-53/1 PROTEIN

Page 1



Tuesday, 18 November 1997 10:33
fig. Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9

Page 2

GCACGGCGAACGGGCCACTACTCCACACCATGCCATGCGCAGCCCCAGCAAGCTCAGCCATATC TCCCGCCTGGAGCTGGTCGAATCCCTGGACTCG 700
CGTGCCGCTTGCCCGGGTGATGAGGGTGTGGTACGGGTACGCGTCGGGGTCGTTCGAGTCGGTATAGAGGGCGGACCTCGACCAGCTTAGGGACCTGAGC
pHH14-3
ORF (1-573bp) = pLM7 ORF
full available ORF HU-Unc53/1 = pLM1 OR
H G E R A H Y S H T M P M R S P S K L S H I S R L E L V E S L D S
GATGAGGTGGACCTCAAGTCCGGCTACATGAGCGACAGTGACCTCATGGGCAAGACCATGACGGAGGATGATGACATCAC TACCGGCTGGGATGAAAGCA 800
CTACTCCACCTGGAGTTCAGGCCGATGTACTCGCTGTCACCTGGAGTACCCGTTCTGGTACTTGCCCTCTACTACTGTAGTGATGGCCGACCTTACTTTCTGT
pHH14-3
ORF (1-573bp) = pLM7 ORF
full available ORF HU-Unc53/1 = pLM1 OR
D E V D L K S G Y M S D S D L M G K T M T E D D D I T T G V D E S
GCTCCATCAGTAGTGGACTCAGCGATGCCTCAGACAATCTCAGTTCAGAAGAATTCATGCCAGCTCCTCACTCAACTCCCTCCCAAGTACTCCCACTGC 800
CGAGGTAGTCATCACC TGAGTCGCTACGGAGTCTGT TAGAGTCAAGTCTTCTTAAGTTACGGTCGAGGAGTGAGTTGAGGGAGGGTTTCATGAGGGTGACG
pHH14-3
pCB212
ORF (1-579bp) = pLM7 ORF
full available ORF HU-Unc53/1 = pLM1 OR
S S I S S G L S D A S D N L S S E E F N A S S S L N S L P S T P T A
TTCTCGCAGGAACCTCAACAATAGTGCTACGCACAGACTCAGAGAAGCGCTCACTGGCAGAAAGTGGGCTGAGCTGGTTTAGTGAATCAGAGGAGAAAGCC 900
AAGAGCGTCCCTTGASTTGTATCAGGATGCGTGTCTGAGTCTCTCGCGAGTGACCGTCTTACCCGACTCGACCAATCAGTCTAGTCTCCCTCTTTCTGG
pHH14-3
pCB212
full available ORF HU-Unc53/1 = pLM1 OR
S R R N S T I V L R T C S E K R S L A E S G L S V F S E S E E V A
CCTAAAAAATGGAGTACGACAGTGGTAGCCTGAAGA TGAACCTGGGACTTCTAAGTGGCGGAGGAGCGGCTGAGAGCTGTGATGATTTCACCAAGG 1000
GGATTTTITGACCTCATGCTGTACCATCGGACTTCTACCTTGGACCTTGAAGATTCACCGCTCCCTCGCCGGACTCTCGACACTACTAAGTAGGTTCT
pHH14-3
pCB212
full available ORF HU-Unc53/1 = pLM1 OR
L E L D S G S L V E P G T S K V R R E P E S C D D S S I

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Fig 9

Page 3

fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

GTGGAGAACTGAAAAAGCCATCAGCCTGGGCCACCTGGTTCCTGAAGAAGGGCAAGACCCACCTGTGGCTGTAACCTCCCCCATCACTCACACAGC
CCTCTTGACTTTTTCGGGTAGTCGGACCCGGTGGGACCAAGGGACTCTTCCCGTTCTGGGGTGGACACCGACATTGAAGGGGGTAGTGAGTGTGCG

pHH14-3

pCB212

full available ORF HU-Unc53/1 = pLM1 OR

G G E L K K P I S L G H P G S L K K G K T P P V A V T S P I T H T A

CCAGAGTGGCCTCAAAGTCGCAGGCAAACTGAGGGCAAGCTTACAGACAAGGGTAAGCTTGCAAGTGAAGAATCTGGGCCTCAACGCCTCTCTCTGAT
GGTCTCACGGGAGTTTCAGCGTCCGTTTGACCTCCCTTCGATGTCTGTTCCCATTCGAACGTCACCTCTTATGACCCGAGGTTGCGAGGAGGAGACTA

pHH14-3

pCB212

full available ORF HU-Unc53/1 = pLM1 OR

Q S A L K V A G K P E G K A T D K G K L A V K N T G L Q R S S S D

GCTGGTCGGGACCGCTGAGTGATGCTAAGAAGCCCCCTCGGGCATTGCTCGCCCCCTCCACTTCGGGATCCTTTGGCTACAAGAAGCCTCCTCTGCGCA
CGACCAGCCCTGGCGGACTCACTACGATCTTTCGGGGGGAGCCCGTAACGAGCGGGGAGGTGAAGCCCTAGGAAACCGATGTTCTTCGGAGGAGGACGGT

pHH14-3

pCB212

full available ORF HU-Unc53/1 = pLM1 OR

A G R D R L S D A K K P P S G I A R P S T S G S F G Y K K P P P A

CAGGCACAGCCACTGTGATGCAAACTGGTGGTTCAGGCACTCTCAGCAAGATCCAGAAGTCTCAGGCATCCCTGTCAAGCCAGTAAATGGCGCAAGAC
GTCTGTGCGGTGACAGTACGTTTGACCAACCAAGTCGTTGAGAGTCGTTTAGGTCTTCAGGAGTCCGTAGGGACAGTTTCGGTCAATTACCCCGGTTCTG

pHH14-3

pCB212

full available ORF HU-Unc53/1 = pLM1 OR

T G T A T V M Q T G G S A T L S K I Q K S S G I P V K P V N G R T T

TAGCTTAGATGTTTCCAACAGTSCAGAGCCAGGATTCCTGGCTCCTGGAGCCCGTTCTAACAATCAGTACCGCAGCCTGCCCGGCCAGCCAAGTCAAGT
ATCGAATCTACAAAGTTGTCAGTCTCGGTCCTAAGTACCGAGGACCTCGGGCAAGATTGTAGGTGATGTCGTCGGACGGGGCCGGTCGGTTCAGTTCA

pHH14-3

pCB212

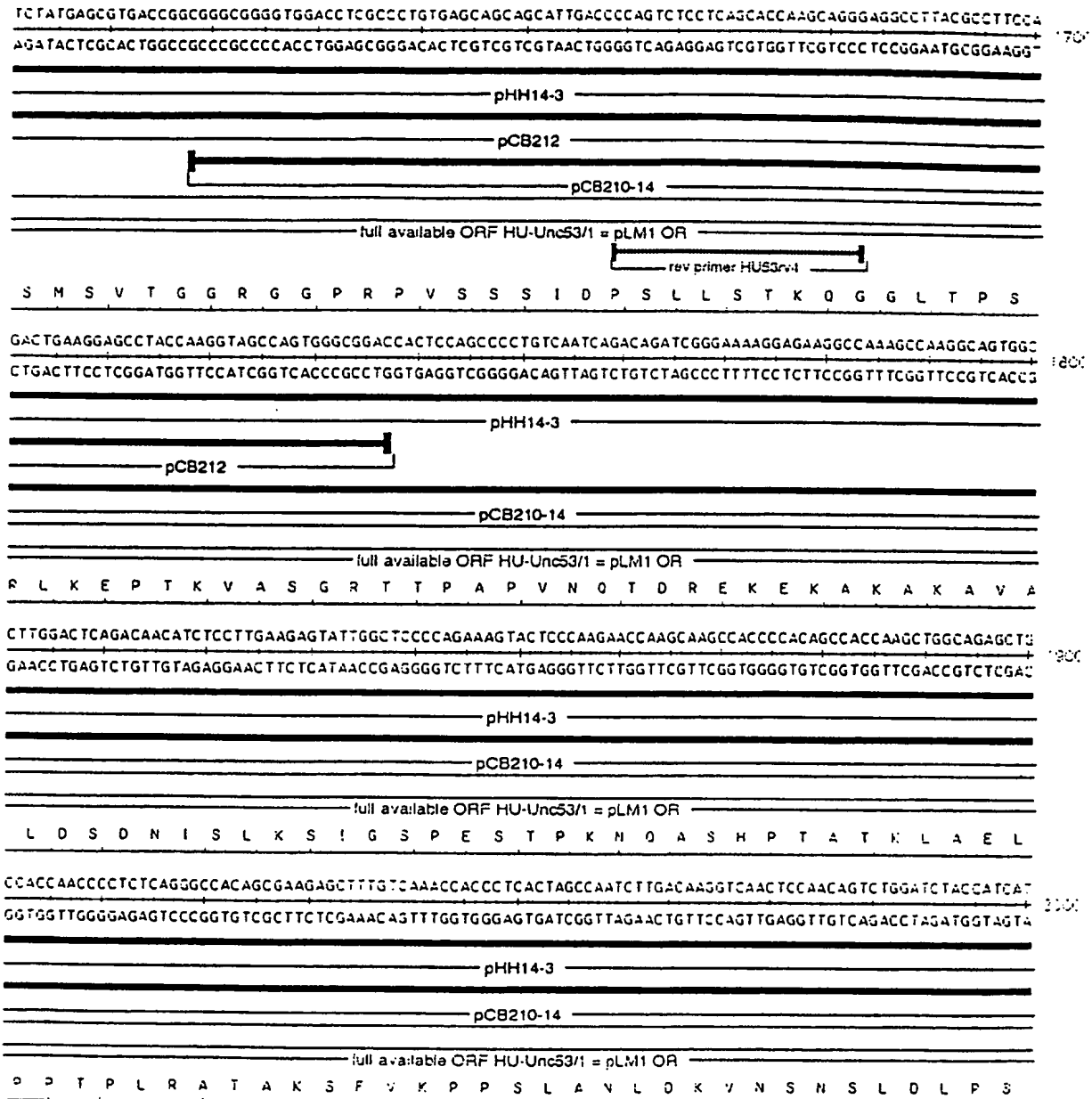
full available ORF HU-Unc53/1 = pLM1 OR

I L D V S H S A E P G F L A P G A R S N I O Y R S L P R P A S S

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fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9

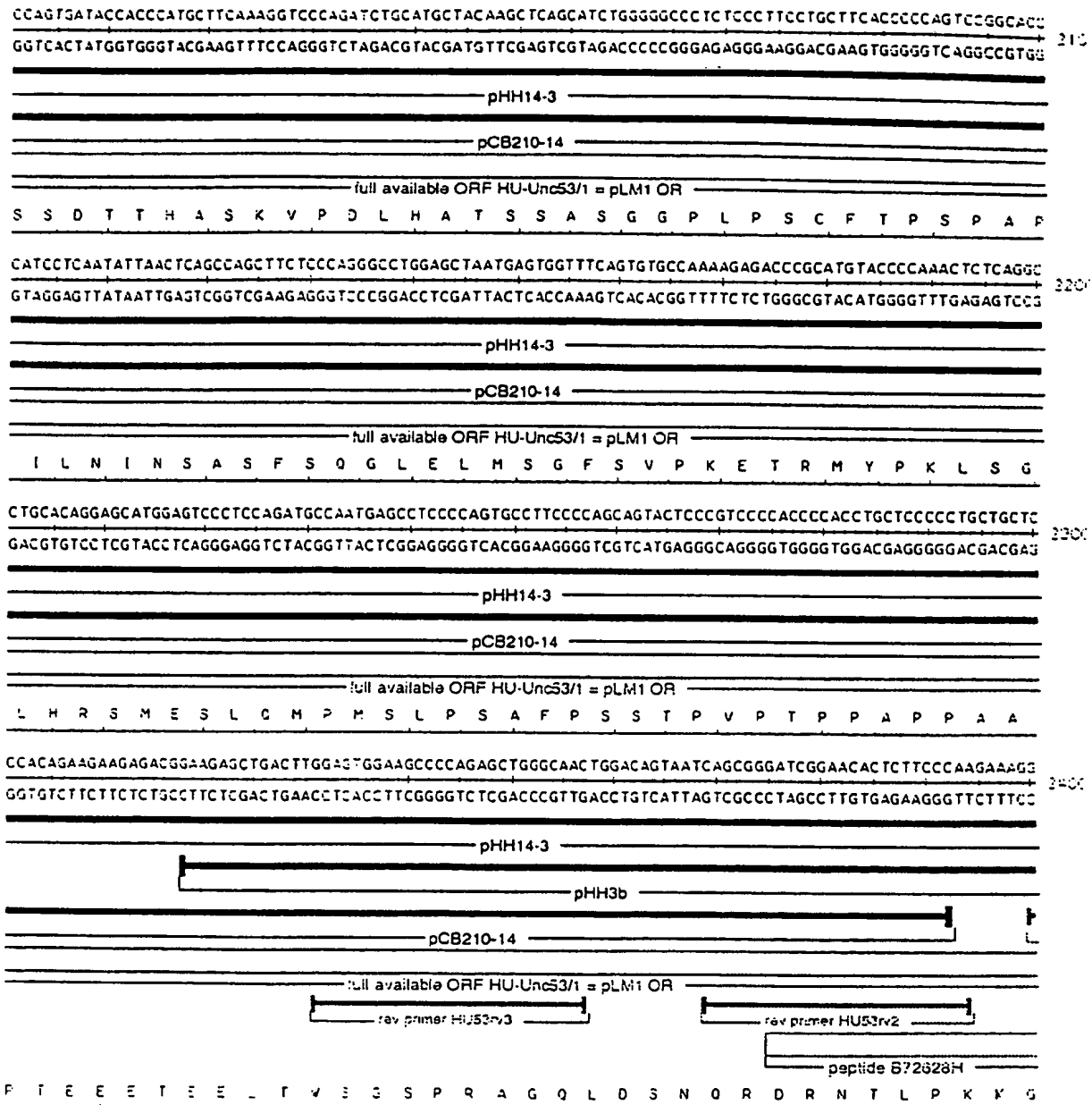
Page 4



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fig. HU-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9

Page 5



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fig HU-Unc53/1 seq (1 > 6013) Site and Sequence

Page 6

GC TCAGGTACCAGCTTCAGTCCCAGGAGGAGACCAAGGAGAGGGGACATTCCCATACCATTTGGTGGGCTGCCTGAATCCGATGACCAGTCAGAGCTGCCT
CGAGTCCATGGTCTGAAGTCAGGGTCTCTCTCTGGTTCTCTCCGCTGTAAGGGTATGGTAACCAACCCGACGGACTTAGGCTACTGGTCAGTCTCGACGGAA 250

pHH14-3

pHH3b

rev primer HU53rv1

full available ORF HU-Unc53/1 = pLM1 OR

L R Y Q L Q S Q E E T K E R R H S H T I G G L P E S D D Q S E L P

TCTCCCCCTGCACCTTCCCATGTCTCTGAGTGCAAGGGGCCAACTTACCAACATAGTGAGTCCCCTGCGGCCACCAAGCAAGCAATCAGCCGCTCCAACT
AGAGGGGGACGTGAAGGGTACAGAGACTCACGTTTCCCGTTGAATGGTTGATCACTCAGGGTGACCGGGTGGTGGGTTCTTAGTGGGCGAGGTTGT 260

pHH14-3

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S P P A L P M S L S A K G Q L T N I V S P T A A T T P R I T R S H

GCATCCCCACCCACGAGGGCGGCTTCGAGCTGTACAGCGGCTCCCAAAATGGGGAGCACCCGTGTCCTGGCCGAGAGACCCAAGGGAATGATTGGGTCAAG
CGTAGGGGTGGGTGCTCCGCGGAAGCTCGACATGTCGCCGAGGGTTACCCCTCGTGGGACAGGGACCGGCTCTCTGGGTTCCCTTAC TAAGCCAGTCC 270

pHH14-3

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S I P T H E A A F E L Y S G S Q M G S T L S L A E R P K G M I R S G

ATCCTTCGAGACCCACGGACGATGTTACGGGCTCAGTCTCTCTCCCTGGCCTCCAGTGCTTCTCCACCTACTCCTCAGCTGAGGAGAGGATGCAATCT
TAGGAAGGCTCTGGGGTGCTGCTACAAGTGCCGAGTACGACAGGGACCGGAGGTACGGAGGAGGTGATGAGGAGTCGACTCCTCTCTCTACGTTAGA 280

pHH14-3

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S F R D P T D D V H G S V L S L A S S A S S T Y S S A E E R M Q S

GACCAATCCGGAAGCTTCGTAGGGAAC TGGAAATCCAGGAAAAAGTGGCCACCTTGACGTCTCAGCTTTCTGCCAATGCTAATCTGGTGGCTGCTT
CTCGTTTAGGCTTCGAAGCATCCCTTGACCTTAGTAGGGTCTTTTACCAGGTGGAAC TGCAGAGTCGAAAGACGGTTACGATTAGACCACCGACGAA 290

U2 ORF = pC8351 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S I R K L R P E L E S S G E K V A T L T S C L S A N A N L V A A

Fig 9

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fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Page 7

TTGAGCAGAGCC TGGTGAATATGACATCCCGCCTGCGACACC TGGCAGAGACGGCCGAGGAGAAGGACACTGAGCTGCTGGATTGCGAGAAACCATAGA
AACTCGTCTCGGACCAC TTATAC TGTAGGGCGGACGCTG TGGACCGCTCTGCGCGGCTCCTC TTCTGTGACTCGACGACCTAAACGCTCTTGGTATCT 300

U2 ORF = pCB251 ORF

PHH3b

full available ORF HU-Unc53/1 = pLM1 OR

F E Q S L V N M T S R L R H L A E T A E E K D T E L L D L R E T I D

CTTCTGAAGAAAAAGAACTCTGAGGCCAGGCGAGTCATTCAGGGAGCCCTTAATGCC TCAGAAACCACACCCAAAGAACTTCGGATCAAGAGACAAAAC
GAAAGACTTCTTTTCTTGAGACTCCGGGTCCGTCAGTAAGTCCCTCGGGAATTACGGAGTCTTTGGTGTGGGTTCTTGAAGCC TAGTCTCTGTTTTG 310

U2 ORF = pCB251 ORF

PHH3b

full available ORF HU-Unc53/1 = pLM1 OR

F L K K K N S E A Q A V I Q G A L N A S E T T P K E L R I K R Q N

TCCTCAGATAGCATCTCAAGCCTCAACAGCATCACTAGCCATTCAGCATCGGCAGCAGCAAGGATGCTGATGCGAAAAAGAAAAAGAGTTGGG
AGGAGTCTATCGTAGAGTTCGGAGTTGTCGTAGTGATCGGTAAGGTCGTAGCCGTCGTCGTTCC TACGACTACGCTTTTCTCTTTTTTCTCAACCC 320

U2 ORF = pCB251 ORF

PHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S S D S I S S L N S I T S H S S I G S S K D A D A K K K K K S V

TCTATGAGCTTCGAAGTTCTTCAACAAAGCGTTCAATATAAAAAAGGGGCCAAGTCAGCTTCCTCATACTCGGATATAGAGGAGATTGCTACACCCGA
AGATACTCGAAGCTTCAAGGAAGTTGTTTCGCAAGTCATATTTTCCCCGGGTTTCAGTCGAAGGAGTATGAGCCTATATCTCCTCTAACGATGTGGGCT 330

U2 ORF = pCB251 ORF

PHH3b

full available ORF HU-Unc53/1 = pLM1 OR

V Y E L R S S F N K A F S I K K G P K S A S S Y S D I E E I A T P D

CTCTCAGCCCCCTCATCCCCAAACTACAGCATGGTCTACAGAGACTGCTTCACCCCTCCATCAAGTCTCTCCACCTTGTCTCCGTGGGCACTGATGTC
GAGAACTCGGGGGAGTAGGGGTTTGATGTCTGTACCAAGATGCTCTGACGAAGTGGGAGGTAGTTCAAGAGGTGGAAACAGGAGGCCACCCCTGACTACAG 340

U2 ORF = pCB251 ORF

PHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S S A P S S P K L Q F G S T E T A S P S I K S S T L S S V S T D V

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Fig 9

Page 5

HU-Unc53/1 seq (1 > 6013) Site and Sequence

ACCC .GCCCTGCTACCCAGCCCCACACTAGGCTGTTCATGCAATGAGGAGGAGGCCAGAGAAGAAGGAGGTATCGGAGCTGCGCTCTGAGC 3500
TGGCTCCCGGGACGAGTGGGTTCGGGGGTGTGATCCSACAAGGTACGTTACTCTCTCTCTCGGTCTCTTCTTCTTCCATAGCTTCGACGCGAGACTCG

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

T E G P A H P A P H T R L F H A N E E E E P E K K E V S E L R S E

TATGGGAGAAGGAAATGAAGCTTACAGACATCCGCTTGGAGGCCCTCAACTCTGCCACCAACTGGATCAGCTTCGGGAGACCATGCACAAATGCAGTT 3600
ATACCTCTTCTCTTACTTCGAATGTCTGTAGGCGAACCTCCGGGAGTTGAGACGGGTGGTTGACCTAGTCGAAGCCCTCTGGTACGTGTTGTACGTCAA

U2 ORF = pCB251 ORF

pHH3b

peptide B72627H

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

L V E K E M K L T D I R L E A L N S A H Q L D Q L R E T M H N M Q L

GGAGGTGGACCTGCTGAAAGCAGAGAATGACCGACTGAAGGTAGCCCCAGGCCCTCATCAGGCTCCACTCCAGGGCAGGTCCCTGGATCATCTGCATTA 3700
CCTCCACCTGGACGACTTTCGTCCTTACTGGCTGACTTCCATCGGGGTCCGGGGAGTAGTCGAGGTGAGGTCCCGTCCAGGGACCTAGTAGACGTAA

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

E V D L L K A E N D R L K V A P G P S S G S T P G Q V P G S S A L

TCTTCCCCACGCCCTCTCCCTAGGCCCTGSCACTCACCATTCTTCCGCCCCAGTCTTGACAGACACAGACCTGTCAACCATGGATGGCATCAGTACTTGTG 3800
AGAAGGGGTGCGGCGAGGGATCCGGACCGTGAGTGGGTAAAGGAAGCCGGGTGAGAAGCTGTGTGCTGGACAGTGGGTACCTACCGTAGTCATGAACAC

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

S S P R R S L G L A L T H S F G P S L A C T D L S P M D G I S T C

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fig. 9 HU-Unc53/1 seq (1 > 6013) Site and Sequence

Page 9

GTCCAAAGGAGGAAGTGACCCCTCCGGGTGGTGGTGAGGATGCCCCCGCAGCACATCATCAAAGGGGACTTGAAGCAGCAGGAATTCTTCTGGGCTGTAG
CAGGTTTCTCTCTTCACTGGGAGGCCACCAACCACTCTACGGGGGCGTCGTGTAGTAGTTTCCCTGAACCTCGTCGTCTCTAAGAAGGACCCGACATC

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

G P K E E V T L R V V V R M P P O H I I K G D L K Q Q E F F L G C S

CAAGGTCAGTGGAAAAGTTGACTGGAAGATGCTGGATGAAGCTGTTTTCAGGTGTTCAAGGACTATATTCTAAATGGACCCAGCCTCTACCTGGGA
GTTCAGTCACCTTTTCAACTGACCTTCTACGACCTACTTCGACAAAAGGTTCAACAAGTTCCTGATATAAAGATTTTACCTGGGTGGGAGATGGGACCT

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

K V S G K V D V K M L D E A V F Q V F K D Y I S K M O P A S T L G

CTAAGCACTGAGTCCATCCATGGCTACAGCATCAGCCAGCTGAAACGAGTGTGGATGCAGAGCCCCCGAGATGCCTCTTGGCGTCGAGGTGTCAATA
GATTCGTGACTCAGGTAGGTACCGATGTCGTAGTCGCTGACCTTTGCTCACAACCTACGCTCTCGGGGGGCTCTACGGAGGAACGGCAGCTCCACAGTTAT

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

L S T E S I H G Y S I S H V K R V L D A E P P E M P P C R R G V N

AGATATCAGTCTCCCTCAAAGGCTCTGAAGGAGAAATGGCTCGACAGCCTGCTGTTTCGAGAGCGTGAATCCCAAGCCGATGATGCAGCACTACATAAGGCT
TGATATAGTCAGAGGGAGTTTCCAGACTTCTCTTTTACGACGCTGTCGGACCAACAAGCTCTCGGACTAGGGGTTTCGGCTACTACGTCGTGATGTAATCGGA

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

H I S V S L Y G L E E C V U S L V F E T L I P A P M M Q H I S L

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Fig. 9 HU-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9

Page 16

CCCTCGAGGACCGGGCGCCTCGTCCCTCGGGCCCCAGCGGCACGGGCAAGACCTACCTGACCAATCGCTTGGCCGAGTACCTGGTGGAGCGCTCTGGG
GGACGACTTCGTGGCCGCGGAGCAGGAGAGCCCCGGGTCGCCGTGCCCGTCTTGGATGGACTGGTTAGCGAACC GGCTCATGGACCACCTCGCGAGACCG

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

L L K H R R L V L S G P S G T G K T Y L T N R L A E Y L V E R S G

CGTGAGGTACAGAGGGCATCGTCAGCACCTTCAACATGCACCAGCAGTCTTGCAAGGATCTGCAACTGTATCTTTCCAACCTAGCCAACAGATAGACC
GCACCTCCAGTGTCTCCCGTAGCAGTCGTGGAAGTTGTACGTGGTCGTCAGAACGTTCTTAGACGTTGACATAGAAAGGTTGGATCGGTTGGTCTATCTGG

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

P E V T E S I V S T F N M H Q Q S C K D L Q L Y L S N L A H Q I D

GGGAAACAGGAATTGGGGAATGTCGCCCTGGTGGTCTTATTTGGATGACCTGAGTGAAGCAGGC TCCATCAGTGAGTTGGTCAATGGGGCCCTCACCTGCA
CGCTTTGTCTTTAACCCCTACACGGGGACCACTAAGATAACCTACTGGACTCACTTCGTCGGAGGTAGTCACTCAACCACTTACCCCGGGAGTGGAGCT

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

P E T G I G D V P L A L L D D L S E A G S S E L V N G A L T C I

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fig. 4u-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9

Page 11

G T A T C A . A A A T G T C C C T A T A T T A T A G G T A C C A C C A A T C A G C C T G T A A A A A T G A C A C C C A A C C A T G G C T T G C A C T T G A G C T T C A G G A T G T T G A C C T T C T C C
C A T A G T A T T T A C A G G A T A T A A T A T C C A T G G T G G T T A S T C G G A C A T T T T A C T G T G G G T T G G T A C C G A A C G T G A A C T C G A A G T C C T A C A A C T G G A A G A G G

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

peptide B72629H

Y H K C P Y I I G T T N Q P V K M T P N H G L H L S F R M L T F S

A A C A A C G T G G A G C C A G C C A A T G G C T T C C T G G T T C G T T A C T G A G G A G G A A G C T G G T A G A G T C A G A C A G C G A C A T C A A T G C C A A C A A G G A A G A G C T G C T T C
T T G T T G C A C C T C G G T C G G T A C C G A A G G A C C A A G C A A T G G A C T C C T C C T T C G A C C A T C T A G T C T G T C G C T G T A G T T A C G G T T G T T C C T T C T C G A C G A A G

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

Y N V E P A N G F L V R Y L R R K L V E S D S D I N A N K E E L L

G G S T G C T C G A C T G G S T A C C C A A G C T G T G G T A T C A T C T C C A C A C C T T C C T T G A G A A G C A G C A C C T C A G A C T T C C T C A T C G G C C C T T G C T T C T T T C T G T C
C C C A C G A C T G A C C C A T G G S T T C G A C A C C A T A G T A G A G G T G T G G A A G G A A C T C T T C G T G T C T G G A G T C T G A A G G A G T A G C C G G G A A C G A A G A A G A C A G

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

R V L D V V F K L V Y - L H T F L E K H S T S D F L I G P C F F L S

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fig. 1J-Unc53/1 seq (1 > 6013) Site and Sequence

Fig.

Page 12

GTGTCCCATTTGGCATTGAGGACTTCCGGACCTGGTTTCATTGACCTGTGGAACAACCTCTATCATTTCCCTATCTACAGGAAGGAGCCAAAGGATGGGATAAAG
CAGAGGGTAAGCGTAACCTCTGAAGGCCCTGGACCAAGTAACCTGGACACCTTGTGTGAGATAGTAAGGGATAGATGTCCCTCCCTCGGTTCCCTACCTATTTCT

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

C P I G I E D F R T V F I D L V N N S I I P Y L Q E G A K D G I I

GTCCATGGACAGAAAGCTGCTTGGGAGGACCCAGTGGAAATGGGTCCGGGACACACTTCCCTGGCCATCAGCCCAACAAGACCAATCAAAGCTGTACCAAC
CAGGTACCTGTCTTTTCGACGAACCTCTCTGGGTCACCTTACCAGGCCCTGTGTGAAGGGACCGGTAGTCGGGTTGTTCTGGTTAGTTTCGACATGGTGG

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

V F G C K A A V E D P V E V V R D T L P V P S A Q Q D C S X L Y H

TNCCGCCACCCACCGTGGGCCCTCACAGCATTGCCTCACCTCCCGAGGATAGGACAGTCAAAGACAGCACCCCAAGTTCTCTGGACTCAGATCCCTCTGAT
ACGGGGGTGGGTGGCACCCGGGAGTGTCTTAACGGASTGGAGGGCTCTATCTGTGAGTTTCTGTGCTGGGGTTCAAGAGACCTGAGTCTAGGAGACTA

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

L P P P T V G P H S I A S P P E O R T V K O S T P S S L D S C P L I

Tuesday, 18 November 1997 10:34
file -lu-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9

Page i 3

GGCCATGCTGCTGAAACCTTCAAGAAGCTGCCAACTTATCTGAGTCTCCAGATCGAGAACCCTCTGGACCCCAACCTTCAGGCAACACTTTAAGGCTTC
CTGGTACGACGACTTTGAAGTCTTCGACGGTGTATGTAATCAGAGGCTTAGCTCTTTGGTAGGACCTGGGGTTGGAAGTCCGTGTGAAATTTCCCAAG

U2 ORF = pCB251 ORF

- pHH3b

- U4 ORF = pCB201 ORF

- full available ORF HU-Unc53/1 = pLM1 CR

- U3 ORF = pLM5 ORF

- pHH15

- pupilo 372825H

A M L L K L Q E A A N Y I E S P D R E T I L D P N L Q A T L . G F

GGCAATCACTGTGTCACCCCGGACAGCAGAACGCTGGCATCAGCTATCTTAGCTCCTCCTCTCCCTCTCCTCTTTCAGAGCACTGGCTCTCCAGCCCCA
CCGTTAGTGACAGTGGGGGCCTGTCGCTCTTGCACCC TAGTCGATAGAATCGAGGAGGAGAGGGGAGAGGAGAGAAAGTCTCTGTACCCGAGAGGTCGGGGT

- pHH3b

- pHH15

U N H C H P R T A E R W H Q L S . L L L S P L L F Q S T G S P A P

[illegible]

- pHH3b

pH 15

U U E O E G G G D E R G G T G S W C C T F E H F L G R N G G V A F G

GAACTTGTCGCCCCTAAACACATTTACTGGCCCTCTCTAATGACTTTGGGGAAAAGATGATCTGGGCTTTCCCTTGACTTCTTGTTTCAATTACAAAC
 CTGGAACACGGGGGATTTGTGTAATGACCGGAGAGATTACTGAAACCCCTTTTCTACTAAGACCCAGAAAGGGGAACGGAAGAACAAAGTTAATGTTT

- pHH3b

- pHH15

M L C P L N T F T G L L . . L W G X D O S G S F P . L L V S I T M

TCCTGGGCTTTCTGGGGAGGGTTTCAGAAAAACA¹CTAAACAC TGCAGC AG TTCCTAAATGAT TCTCACAAGCAACCCCTGAGAGAGACAG TCTTCTGAGGG
AGGACCCCGAAAGACCCCTCCCAAGTCTTTTGTACTTTTGTGACGTCGTCAGGATTACTTAAGAGTGTTCGTTGGGACTCTCTCTGTCTAGAACACTCC

- pHH3b

- pHH15

W A F W G G V G D T I . H C S S S . M I . S V P E R O S L V R

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fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9

Page 14

AGATCTGGGGGAGGAGGAGGAGCTCCTCAGATTTTCTACAGACCCCTTCCCAATTCCATCACCAC TGCCAACAATCTCTCCCCAGAGATCTGGCTGGAGC
TCTAGACCCCTCCGTCCTTCGAGGAGTCTAAAAGAGTGCTGGGAAGGGTTAAGGTAGTGGTGACGGTTGTTGAGGAGGGGGTCTCTAGACCGACCTCG 570

pHH15

E I V G R Q E A P Q I F S Q T L P N S I T T A N N S S P R D L A G A

CCAGAAAAAGAAGCATGTGGTTTAAAAATGTTTAAATCAATCTGTAAAAGGTAAAAATGAAAAACAAAAACAAGCAACAACAAAAACAATGGAAA 580
GGTCTTTTCTTCGTACACCAAAATTTTACAAATTTAGTTAGACATTTTCCATTTTACTTTTGTTTTGTTTCGTTTGTGTTTTGTTTACCTTT
Q K K K H V V . K M F K S I C K R . K . K N K N K Q T N K K Q V I

AGATGAAGCTGGAGAGAGAGGAACCAAGTTGCCAAGGTAGAGAGCTGCCCGCTCCTGCCCTCTGGATGACATAGGGGACATCAACAAGACGGCTGCCAAC 590
TCTACTTCGACCTCTCTCTCCTTGGTCAACGGTTCCATCTCTCGACGGGCGAGGACGGGAGACCTACGTATCCCTGTAGTTGTTCTGCCGACGGTTGG
R . S V R E R N Q L P R . R A A R S C P L D D I G D I N K T A A H

TGAGAAGTCACCAAAACCACAAAAATAACCTTACAGCC TTCAGGGAAAGACTACCAGCTCTGTCTTCTACCCCTCTAATTTAACAATGCACCGGAATTCAG 600
ACTCTTCAGTGGTTTGGTGTGTTTTATTGGAATGTCGGAAGTCCCTTCTGATGGTCGAGACAGAAAGATGGGAGATTAAATTGTTACGTGGCCTTAAGTC

linker?

L R S H Q T T K I T L C P S G K D Y Q L C L S T L . F N N A P E F S

CTTGGACTTAACC 6013
GAACCTGAATTGG

linker?

L D L T

SEQ ID NO 2: Amino Acid Sequence of Hn-uuc-53/2 Protein

[illegible][illegible]

FD 302 (Rev. 11-27-70)

1288
1289
1290
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1292
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[illegible]

[illegible]

Seq. ID No 3

Nucleic Acid Sequence of
hu-unc-53/1 gene

Page 1

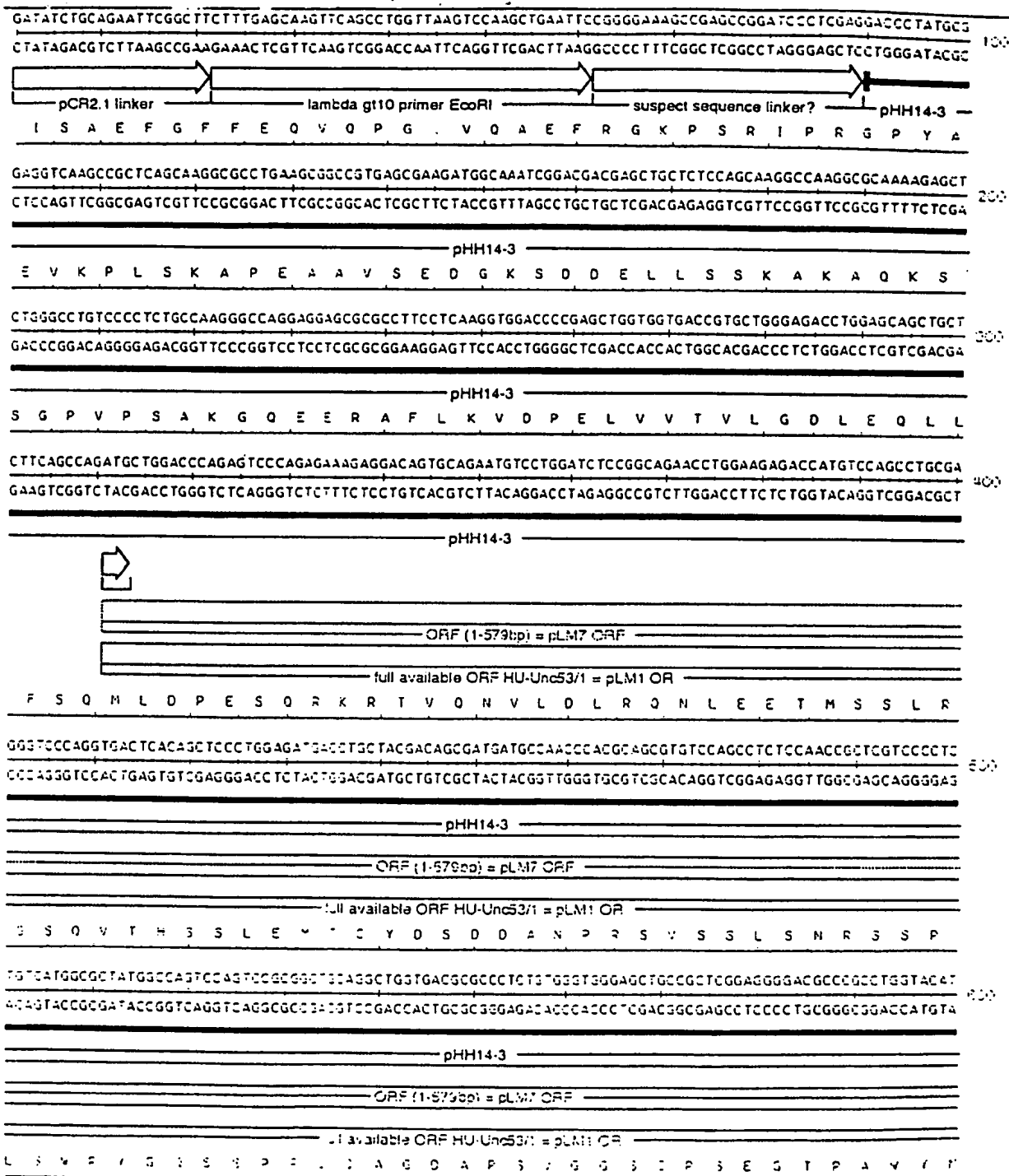


Fig 9

Page 2

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fig Hu-Unc53/1 seq (1 > 6013) Site and sequence

GCACGGCGAACGGGCCCACTACTCCACACCATGCCATGCGCAGCCCCAGCAAGCTCAGCCATATC TCCCGCCTGGAGCTGGTCGAATCCCTGGAATCTG
CGTCCCGCTTGCCCGGGTGATGAGGGTG TGGTACGGGTACGCGCTCGGGGTCTGTCGAGTCGGTATAGAGGGCGGACCTCGACCAGCTTAGGGACCTGAGC
pHH14-3
ORF (1-579bp) = pLM7 ORF
full available ORF HU-Unc53/1 = pLM1 OR
H G E R A H Y S H T M P M R S P S K L S H I S R L E L V E S L D S
GATGAGGTGGACCTCAAGTCCGGCTACATGAGCGACAGTGACCTCATGGGCAAGACCATGACGGAGGATGATGACATCACTACCGGCTGGGATGAAGCA
CTACTCCACCTGGAGTTCAGGCCGATGTACTCGCTGTCACTGGAGTACCCGTTCTGGTACTGCCTCCTACTACTGTAGTGATGGCCGACCTTACTTCTGT
pHH14-3
ORF (1-579bp) = pLM7 ORF
full available ORF HU-Unc53/1 = pLM1 OR
D E V D L K S G Y M S D S D L M G K T M T E D D D I T T G V D E S
GCTCCATCAGTAGTGGACTCAGCGATGCCCTCAGACAATCTCAGTTCAGAAGAATTCAATGCCAGCTCCTCACTCAACTCCCTCCCAAGTACTCCCACTGC
CGAGGTAGTCATCACCTGAGTCGCTACGGAGTCTGTTAGAGTCAAGTCTTCTTAAGTTACGGTCGAGGAGTGAGTTGAGGGAGGGTTCATGAGGGTGACG
pHH14-3
pCB212
ORF (1-579bp) = pLM7 ORF
full available ORF HU-Unc53/1 = pLM1 OR
S S I S S G L S D A S D N L S S E E F N A S S S L N S L P S T P T A
TCTCGCAGGAACCTCAACAATAGTGTACGGCACAGCTCAGAGAAGCGCTCACTGGCAGAAAGTGGGCTGAGCTGGTTTAGTGAATCAGAGGAGAAAGCC
AAGAGCGTCTTGAGTTGTATCAGGATGCGTGTCTGAGTCTCTCGCGAGTGACCGTCTTTCACCGACTCGACCAATCACTTAGTCTCTCTCTGCG
pHH14-3
pCB212
full available ORF HU-Unc53/1 = pLM1 OR
E R R N S T I V L R T C S E K R S L A E S G L S V F S E S E E I A
CCCAAAAAC TGGAGTACGACAGTGGTACCGTCAAAATGGAACCTGGGACTTCTAAGTGGCGGAGGGAGCGGCTGAGAGCTGTGATGATTGATCCAGG
GGATTTTTCGACCTCATGCTGACCATCGGATTTTACCTTTGGACCTGGAAGATTCACCGCTCCCTCGGCGGACTCTGACACTACTAAGTAGGTTCC
pHH14-3
pCB212
full available ORF HU-Unc53/1 = pLM1 OR
L E T C S C S L N E P G T S K W R P E R P E C C C S S I

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Fig 9

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lig Hu-Unc53/1 seq (1>6013) Site at Sequence

GTGGAGAACTGAAAAAGCCCATCAGCCTGGGCCACCTTGGTTCCCTGAAGAAGGGCAAGACCCACCTGTGGCTGTAACTTCCCCATCACTCACAAGG
CACCCTCTTACTTTTTCGGGTAGTCGGACCCGGTGGGACCAAGGGACTTCTTCCCGTTCTGGGGTGGACACCGACATTGAAGGGGGTAGTGAGTGTGTCT

pHH14-3

pCB212

full available ORF HU-Unc53/1 = pLM1 OR

G G E L K K P I S L G H P G S L K K G K T P P V A V T S P I T H T A

CCAGAGTGCCTCAAAGTCGCAGGCAAACCTGAGGGCAAAGCTACAGACAAGGGTAAGCTTGCAGTGAAGAATACTGGGCTCCAACGCTCCTCCTCTGAT
GGTCTCACGGGAGTTTCAGCGTCCGTTTGGACTCCCTTTTCGATGTCTGTTCCTTCGAACTGACCTTCTTATGACCCGAGGTTGCGAGGAGGAGACTA

pHH14-3

pCB212

full available ORF HU-Unc53/1 = pLM1 OR

D S A L K V A G K P E G K A T D K G K L A V K N T G L Q R S S S D

GCTGGTGGGACCGCTTGAGTGATGCTAAGAAGCCCCCTCGGGCATTCGTCGCCCTCCACTTCGGGATCCTTTGGCTACAAGAAGCCTCCTCCTGCCA
CGACCAGCCCTGGCGGACTCACTACGATCTTTCGGGGAGCCCGTAACGAGCGGGGAGGTGAAGCCCTAGGAAACCGATGTCTTCGGAGGAGGACGGT

pHH14-3

pCB212

full available ORF HU-Unc53/1 = pLM1 OR

A G R D R L S D A K K P P S G I A R P S T S G S F G Y K K P P P A

CAGGCACAGCCACTGTCTATGCAAACTGGTGGTTCAGCCACTCTCAGCAAGATCCAGAAGTCTCAGGCATCCC TGTCAGGCCAGTAAATGGGCGCAAGAC
GTCCGTGTCGGGTGACAGTACGTTTGACCAACCAATCCGTGAGAGTCGTCTAGGTCTTCAGGAGTCCGTAGGGACAGTTCGGTCAATTTACCCGGGTCTCT

pHH14-3

pCB212

full available ORF HU-Unc53/1 = pLM1 OR

T G T A T V M C T G S S A T L S K I O K S S G I P V K P V N S R I T

TAGCTTAGATGTTTCCAACAGTCCAGAGCCCAATTCCTTGGCTCCTGGAGCCCGTCTTAACATCCAGTACCGCAGGCTGCCCGGGCCAGCCCAATCAAGT
ATCGAATCTACAAGAGTGTGACGCTCTGGTCTTAAGGACCGAGGACCTCGGGCAAGATTGTAGGTCAATGCGCTCGGACGGGGCCGGTCCGCTTCAGTTCT

pHH14-3

pCB212

full available ORF HU-Unc53/1 = pLM1 OR

L D V S H S A E P C F L A F G A R S N I Q Y R S L P R F A S

Fig 9

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fig Hu-Unc53/1 seq (1 > 6013) Site a: sequence

TCATGAGCGTGACCGCGGGCGGGGTGACCTCGCCCTGTGAGCAGCAGCATTGACCCCACTCTCTCAGCACCAGGAGGGAGGCTTACGGCTTCCA
AGATACTCGCACTGGCCGCCCGCCCACTTGGAGCGGACACTCGTCTGTCTAAGTGGGGTCAGAGGAGTCGTGGTTCTGTCCCTCCGGAATGCGGAAGG

pHH14-3

pCB212

pCB210-14

full available ORF HU-Unc53/1 = pLM1 OR

rev primer HU53rv1

S M S V T G G R G G P R P V S S S I D P S L L S T K Q G G L T P S

GACTGAAGGAGCCTACCAAGGTAGCCAGTGGGCGGACCACTCCAGCCCCGTCAATCAGACAGATCGGGAAAAGGAGAAGGCCAAAGCCAAGGCAGTGGC
CTGACTTCTCGGATGGTTCCATCGGTACCCGCCCTGGTGAGGTCGGGGACAGTTAGTCTGTCTAGCCCTTTTCTCTTCCGGTTTCGGTTCCGTCACCC

pHH14-3

pCB212

pCB210-14

full available ORF HU-Unc53/1 = pLM1 OR

R L K E P T K V A S G R T T P A P V N Q T D R E K E K A K A K A V A

CTTGGACTCAGACAACATCTCCTTGAAGAGTATTGGCTCCCCAGAAAGTACTCCCAAGAACCAAGCAAGCCACCCACAGCCACCAAGCTGGCAGAGCTG
GAACCTGAGTCTGTTGTAGAGGAACCTTCATAACCAAGGGGCTTTTCATGAGGGTCTTGGTTCTGGTTCGGTGGGGTGTCTGGTGGTTCGACCGTCTCGAC

pHH14-3

pCB210-14

full available ORF HU-Unc53/1 = pLM1 OR

L D S D N I S L K S : G S P E S T P K N Q A S H P T A T I L A E L

CCACCAACCCCTCTCAGGGCCACAGCGAAGAGCTTCTCAAAACACCCCTACTAGCCAATCTTGACAAGGTCAACTCCAAACAGTCTGGATCTACCATCAT
GGTGGTTGGGAGAGTCCCGGTGTCGCTTCTCGAAACAGTTGGTGGGAGTATCGGTTAGAAGTGTTCAGTTGAGGTGTCTAGACCTAGATGGTAGTA

pHH14-3

pCB210-14

full available ORF HU-Unc53/1 = pLM1 OR

P P T P L R A T A K S F V V P P S L A N L D K V N S N S L D L P S

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fig. HU-Unc53/1 seq (1 > 6013) Site. , Sequence

Fig 9

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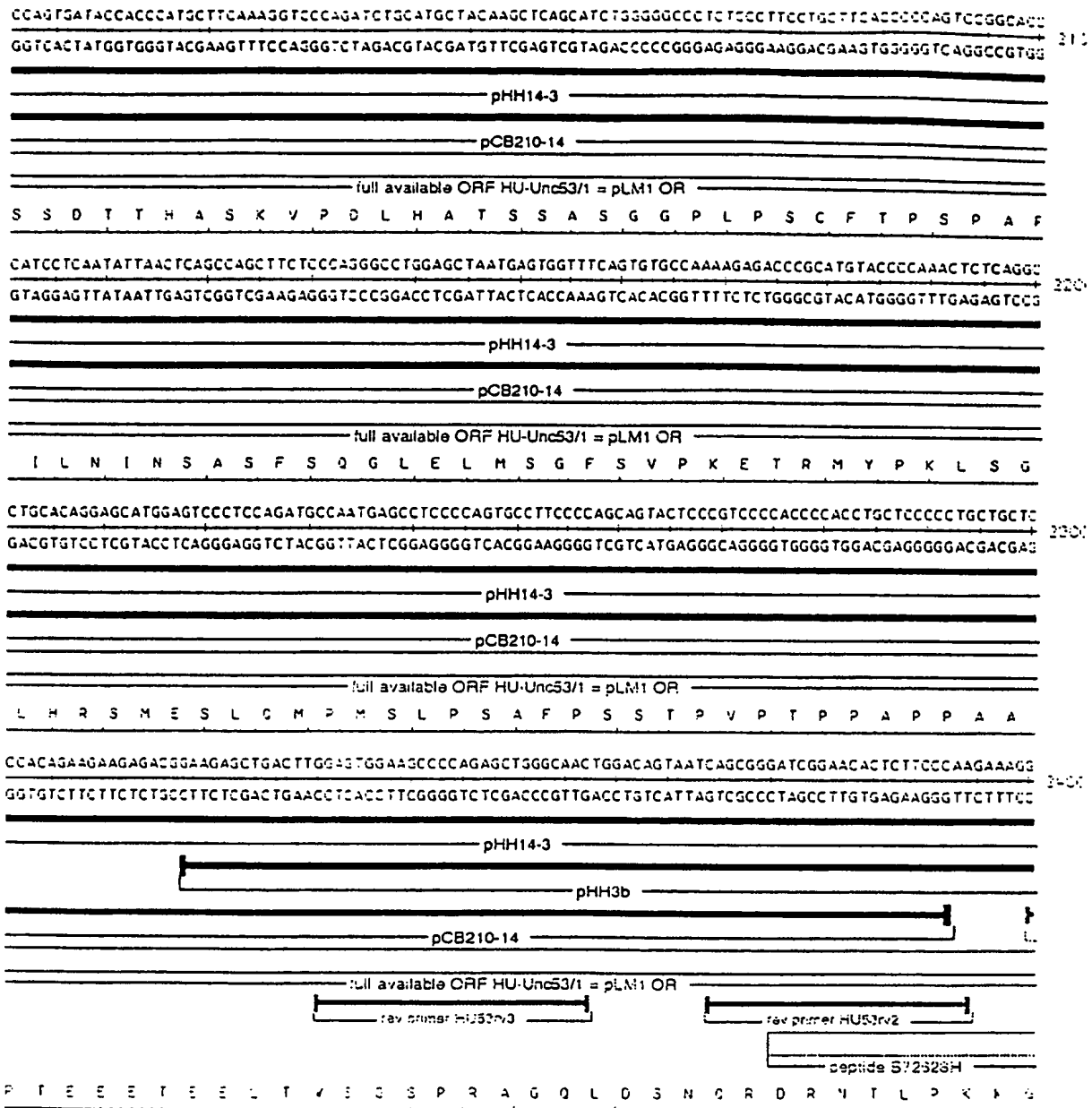


Fig 9

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fig HU-Unc53/1 seq (1 > 6013) Site and sequence

GCTCAGGTACCAGCTTCAGTCCCAGGAGGAGACCAAGSAGAGGGGACATTCCCATACCATTGGTGGGCTGCCTGAATCCGATGACCAGTCAAGAGCTGCCT
CGAGTCCATGGTCGAAGTCAGGGTCCCTCTCTGGTTCCTCTCCGCTGTAAAGGGTATGGTAACCCACCGACGGACTTAGGCTACTGGTCAGTCTCGACGGG 350

pHH14-3

pHH3b

rev primer HU53rv1

full available ORF HU-Unc53/1 = pLM1 OR

L R Y Q L Q S Q E E T K E R R H S H T I G G L P E S D D Q S E L F

TCTCCCCCTGCACCTTCCCATGTCTCTGAGTGCAGAGGCCAACTTACCAACATAGTGAGTCCCAGTGGGCCACCCAGGCCAAGAAATCACCCGCTCCAACA
AGAGGGGGACGTGAAGGGTACAGAGACTCACGTTTCCCGGTGAATGGTTGTATCACTCAGGGTGACGCCGGTGGTGGCGTTCTTAGTGGGCGAGGTTGT 360

pHH14-3

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S P P A L P H S L S A K G Q L T N I V S P T A A T T P R I T R S N

GCATCCCCACCCACGAGGGGGCTTCGAGCTGTACAGTGGCTCCCAAAATGGGGAGCACCTGTCCCTGGCCGAGAGACCCAAGGGAATGATTCGGTCAGG
CGTAGGGGTGGGTGCTCCGCCGAAGCTCGACAATGCGCCGAGGGTTTACCCCTCGTGGGACAGGGACCGGCTCTCTGGGTTCCTTACTAAGCCAGTCC 370

pHH14-3

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S I P T H E A A F E L S G S Q M G S T L S L A E R P K G M I R S G

ATCTTCCGAGACCCACGGACGATGTTACGCTTCACTCTCTCTCCCTGGGCTCCAGTGCCTTCTCCACCTACTCTCAGCTGAGGAGAGGATGCAATCT
TAGGAAGGCTCTGGGGTGCCTGCTACAAGTGGCTAGTCAAGACAGGGACCGGAGGTACGGAGGAGGTGGATGAGGAGTCGACTCCTCTCTCTACGTAGA 380

pHH14-3

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S F R D P T D C V H S S L S L A S S A S S T Y S S A E E R M Q S

GAGCAAAATCCGGAAGCTTCGTAGGGAATGGAAATATCCGAGGAAAAAGTGGCCACCTTGACGTCTCAGCTTTCGCAATGCTAATCTGGTGGCTGCT
CTGGTTTAGGGCTTCGAAGCATCCCTTGACCTTATACGCTCTCTTTCACCGGTGGAACTGCAGAGTCGAAAGACGTTACGATTAGACCACCGAGGAA 390

U2 ORF = pCD351 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S I P K L P D E L E S E N V A T L T S Q L S A N A N L V A A

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Fig 9

fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Page,

TTGAGCAGAGCC TGGTGAATATGACATCCCGCCTGCGACCTGGCAGAGACGGCCGAGGAGAAGGACACTGAGCTGCTGGATTTCGGAGAAACCATAGA
AATCTGCTCGGACCACCTTATACGTAGGGCGGACGCTGTGGACCGTCTCTGCCGGCTCCTCTTCCTGTGACTCGACGACCFAAACCGCTCTTTGGTATCT

300

U2 ORF = pC5251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

F E Q S L V N M T S R L R H L A E T A E E K D T E L L D L R E T I D

CTTCTGAAGAAAAAGAACTCTGAGGCCAGGCAGTCATTCAAGGAGCCCTTAATGCCTCAGAAACCACACCCAAAGAATTCGGATCAAGAGACAAAC
GAAAGACTTCTTTTCTTGAGACTCCGGGTCGGTCATAAGTCCTCGGAATTACGGAGTCTTTGGTGTGGGTTCTTGAAGCCTAGTTCCTCTGTTT

310

U2 ORF = pC8251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

F L K K K N S E A Q A V I O G A L N A S E T T P K E L R I X R Q H

TCCTCAGATAGCATCTCAAGCCTCAACAGCATCAC TAGCCATTCCAGCATCGGCAGCAGCAAGGATGCTGTATGCGAAAAAGAGAAAAAGAGTTGGG
AGGAGTCTATCGTAGAGTTCGGAGTTGTCGTAGTGATCGGTAAGGTCGTAGCCGTCGTCTCC TACGACTACGCTTTTCTCTTTTTTCTCAACCC

320

U2 ORF = pC8251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S S D S I S S L N S ! T S H S S I G S S K D A D A K K K K K S W

TCTATAGCTTCGAAGTTCTCTCAACAAAGCGTTCAGTATAAAAAAGGGGCCAAGTCAGCTTCCTCATACTCGGATATAGAGGAGATTGCTACACCGA
AGATACTCGAAGCTTCAAGGAAGTTGTTTCGCAAGTCATATTTTCCCGGGTTCAGTCGAAGGAGTATGAGCCTATATCTCTCTAAGCATGTGGG

330

U2 ORF = pC8251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

V Y E L R S S F N K A F S I K X G P K S A S S Y S D I E E I A T P D

CTCTTCAGCCCCGTCATCCCCCAACTACAGCATGGTCTACAGAGACTGCTTCACCCCTCCATCAAGTCCCTCCACCTTGTCTCCCTGGGCACTGATGT
GAGAAAGTGGGGGAGTAGGGGTTTGATGCTTACCAAGATGCTCTGACCAAGTGGGAGTAGTTCAGGAGGTGGAAACAGGAGGCCACCCGTGACTACAG

340

U2 ORF = pC8251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S A P S S P K L Q - S S T E T A S P S I X S S T L S S V S T D

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lin HU-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9

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ACCC .GCCCTGCTACCCAGCCCCACACTAGGC TGTTCATGCAAA TGAGGAGGAGGAGCCAGAGAAGAAGGAGGTATCGGAGCTGCGCTCTGAGG
TGGCTCCCGGGACGAGTGGGTCCGGGGGTGTGATCCSACAAGGTACGT TACTCCTCCTCCTCGGTC TC TCTTCTTCCATAGCCTCGACSCGAGACTCG 3500

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

T E G P A H P A P H T R L F H A N E E E E P E K K E V S E L R S E

TATGGGAGAAGGAAATGAAGCTTACAGACATCCGCTTGGAGGCCCTCAACTCTGCCACCAACTGGATCAGCTTCGGGAGACCATGCACAACATGCAGT
ATACCCTCTTCTTTACTTTCGAATGCTGTAGGCGAACC TCCGGGAGT TGAGACGGGTGGTTGACCTAGTCGAAGCCCTCTGGTACGTGTTGTACGTCAA 3600

U2 ORF = pCB251 ORF

pHH3b

peptide B72927H

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

L V E K E M K L T D I R L E A L N S A H Q L D Q L R E T M H N M Q L

GGAGGTGGACCTGCTGAAAGCAGAGAATGACCGACTGAAGGTAGCCCCAGGCCCTCATCAGGCTCCACTCCAGGGCAGGTCCCTGGATCATCTGCATT
CCTCCACCTGGACGACTTTCTGCTCTTACTGGCTGACTTCCATCGGGGTCCGGGGAGTAGTCCGAGGTGAGGTCCCGTCCAGGGACCTAGTAGACGTAA 3700

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

E V D L L K A E N D L K V A P G P S S G S T P G Q V P G S S A L

TCTTCCCGACGCCGCTCCCTAGGCC TGGACTTATCCATTCCTTCGGCCCCAGTCTTGACACACAGACCTGTACCCATGATGGCATCAGTACTTGT
AGAAGGGGTGGCGCGAGGGATCCGGACGCTGATGGGTAAGGAAGCCGGGTCAGAACGCTGTGTCTGGACAGTGGGTACC TACCGTAGTATGAACAG 3800

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

S S P R R S L G L A L - S F G P S L A D T C L S P H D G : S T C

Fig. 9

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fig. 9: Unc53/1 seq (1 > 6013) Site and Sequence

CCCTCTGAAAGACCGGCGCCTCGTCCCTCTCGGGCCCCAGCGGCACGGGCAAGACCTACCTGACCAA*CGCTTGGCCGAGTACC TGGTGGAGCGCTCTGGG
GGACGACTCTCGTGGCCGCGGAGCAGGAGAGCCCGGGGTGCGCGTGCCTGTTCTGGATGGACTGGTTAGCGAACC GGCTCATGGACCACTCTGGGAGACCG 430

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

L L K H R R L V L S G P S G T G K T Y L T N R L A E Y L V E R S G

CGTGAGGTCACAGAGGGCATCGTCAGCACCTTCAACATGCACCAGCAGTCTTGCAAGGATCTGCAAC TGTATCTTTCCAACCTAGCCAAC CAGATAGACC
GCACCTCCAGTGTCTCCCGTAGCAGTCGTGGAAGTTGTACGTGGTCGTGACAACGTTCTTAGACGTTGACATAGAAAGGTTGGATCGGTTGGTCTATCTGG 440

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

P E V T E S I V S T F N M H Q Q S C K D L Q L Y L S N L A N Q I D

GGGAACAGGAATTGGGATGTGCCCCCTGGTGA*TTCTATGGATGACCTGAGTGAAGCAGGCTCCATCAGTGAAGTTGGTCAATGGGGCCCTCACTGCAA
GGTGTGTCCTTAACCCCTACACGGGGACCACTAAGATAACC TACTGGACTCACTTCGTCCGAGGTAGTCACTCAACCACTTACCCCGGGAGTGGAGGT 450

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

P E T S I G C V P L L L C D L S E A G S I S E L V Y G A L T C F

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Fig 9

Page 11

fig. HU-Unc53/1 seq (1>6013) Site Sequence

GTATCA. AAATGTCCCTATATTATAGGTACCAACCAATCAGCCCTGTAAGAAATGACACCCCAACCATGGCTTGCACCTGAGGCTTCAGGATGTTGACCTTCTCC
CATAGTATTACAGGATATAATATCCATGGTGGTTAGTCGGACATTTTACTGTGGGTTGGTACCGAACGTGAACTCGAAGTCTTCAAC TGSAGAGG

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

pupido 872626H

Y H K C P Y I I G T T N Q P V K M T P N H G L H L S F R M L T F S

AACAACGTGGAGCCAGCCAATGGCTTCTGGTTCGTTACCTGAGGAGGAAGCTGGTAGAGTCAGACAGCGACATCAATGCCAACAAGGAAGAGCTGCTTC
TTGTTGCACCTCGGTCGGTTACCGAAGGACCAASCAATGGAC TCCTCCTTCGACCATCTCAGTCTGTGCTGTAGTTACGGTGTTCCTTCGACGAAG

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

N N V E P A N G F L V R Y L R R K L V E S D S D I N A N K E E L L

GGTGGTGGAGTGGGTAACCAAGCTGTGGTATCACTCCACACCTTCTTTSAGAAGCACAGCACCTCAGACTTCTTCATCGGCCCTTGCTTCTTCTGT
CCACGAGGTGACCATGAGGTGACACCATAGTAGAGGTGTGGAAGGAAC TCTTCGTGTCTGAGAGTCTGAAGGAGTACCGGGGAACGAAGAAAGAAAG

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

P V L G A V P K L I I - - P T F L E X H S T S D F L I G P C F F L S

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fig. 9 HU-Unc53/1 seq (1 > 6013) Site an. sequence

Fig. 9

page 12

GTGTGGCATTGGCATTGAGGACTTCCGGACCTGGTTTCATTGACCTGTGGAACAACCTATCATTTCCCTATCTACAGGAGAGGAGCCAGGGATGGGATAAAC
CAGAGGGTAAACCGTAACCTCCTGAAGGCCCTGGACCAAGTAACCTGGACACCTTGTGAGATAGTAAGGGATAGATGTCTCTCTGGTTCCCTACCTATTTCT

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

C P I G I E D F R T V F I D L V N N S I I P Y L Q E G A K Q G I I

GTCCATGGACAGAAAGCTGCTTGGGAGGACCCAGTGGAAATGGGTCCGGGACACACTTCCCTGCECCATCAGCCCAACAAGACCAATCAAAGCTGTACCACT
CAGGTACCTGTCTTTTCGACGAACCTCCTGGGTCACTTACCCAGGCCCTGTGTGAAGGGACCGGTAGTCGGGTGTCTTGGTTASTTTTCGACATGGTGG

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

V - S C K A A V E D P V E V V R D T L P W P S A Q Q D Q S K L Y H

TGCCCCCAGCCACCGTGGGCCCTCAGCATTGCTTCACTCCCCGAGGATAGGACAGTCAAAAGACAGCAGCCCAAGTTCTCTGGACTCAGATCTCTGAT
AGGGGGTGGGTGGCAGCCCGGAGTGTGTAAAGGAGTGGAGGGCTCTATCTGTGAGTTTCTGTGTGGTGGGTTCAAGAGACCTGAGTCTAGGAGACTA

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

L P P F T V G P F S I A E P P E O R T V K D S T P S S L D S C P L I

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| fig | tu-Unc53/1 seq (1 > 6013) | Site a | Sequence |
|-----|---------------------------|--------|----------|
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Fig. 9

Page 13

GGGCATGCTGCTGAAACCTCAAGAAGCTSCCAAC TACAT TGAGTC FCCAGATGGAGAAACCATCTCGGACCCC AACCTTCAGGCAACAC TTTAAGGGTTC
CUGGTACGACGACTTTGAAGTCTCTTGACGGGTGATGTAAC TCAGAGGTC TAGCTCTCTTGGTAGGACCTGGGGT TGGAGTCCGTTGTGAAT TCCCAGG

U2 ORF = 508251 ORF

- pHH3b

U4 ORF = pCB201 ORF

- full available ORF HU-Unc53/1 = pLM1 CR

- U3 ORF = pLM5 ORF

- pHH15

- pasivus B72025H

A M L L K L Q E A A N Y I E S P D R E T I L O P N L Q A T L . G F

GGCAATCACTGTGACCCCCGGACAGCAGAACGCTGGCATCAGCTATCTTAGCTCCTCCTCTCCCTCTCCTCTTTCAGAGCACGGGCTCTCCAGCCCCA3
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- pHH3b

-pHH15

U N H C H P R T A E R W H Q L S . L L L S P L L F Q S T G S P A P

GAGGAGAACAGGAGGAGAGATGAAGAGAGAGGACAGGTCTTGGTGTGTACCTTGGAGAACTTCTAGGAAGGAAATGGTGGGTGGCGTTTG
 CTCTCTGTGTCTGGCTCTCTCTCTACTTTCTCTCTCTCTCCCAAGAACCCAGGACATGGAACTCTTGAAGGATCTCTCTTACCACCCACCCGAACG

- рННЗб

- pHH15

U U E D E S G G D E R S S T G S W C C T F E N F L G R N G G V A F G

GAACTTGTGGCCCCATAACACATTTACTGGCCTCTCTAATGACITTTGGGGAAGAATGATTCTGGGTCTTTCCCTTGACTTCTTTTCAATTACAAA
 CTTCAACACGGGGGATTTGTGTAATGACCGGAGAGATTAATGAAACCCCTTTTCTACTAAGACCCAGAAAGGGAATGAAGAACAAAGTTAAATTGTTT

- pHH3b

- pH 15

N L C P L N T F T G L L . . L V G K O O S S S F F . L L V S I T N

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 AGAACCCGAAAGACCCCTCCCCAAGTCTTTTGTATTTTGTGACGTCGTCGAAGGATTACTTAAGAGTGTTCGTGGGACCTCTCTCTGTGCAACACATCC

~ pHH3b

PHH15

A A F V G O V O A I I H C S O S M I I I S P E E O I Y F

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| 1 | P | E | S | A | N | E | A | P | S | M | S | T | M | G | T | M | A | S | S | S | S | A | V | S |
| 0841 | A | E | S | C | A | C | A | C | A | C | A | C | A | C | A | C | A | C | A | C | A | C | A | C |
| | T | O | S | S | T | O | S | T | O | S | T | O | S | T | O | S | T | O | S | T | O | S | T | O |
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| | T | O | S | | | | | | | | | | | | | | | | | | | | | |

[illegible]

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 -1 D S N S M S M H E D I L S S L S T L O G P G A
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 -1 R P P L L L T A F H L H P P H P E D D F L S Q P P
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 -1 P P P D T E D T S R E I I A V E M K K K K K K
 CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ
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 -1 K K K K
 CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ CTTGZGZGZ
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[illegible]

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SEQUENCE ID NO. 6

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(ig 54 pLM1 (1 > 8285) Site and Sequence

Enzymes: 72 of 148 enzymes (Filtered)

Settings: Circular, Certain Sites Only, Standard Genetic Code

Page 1

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CACCGTGAAAGCCCTTTACACGCGCTTGGGATAACAATAAAAGATTATGTAAGTTTATACATAGGCGAGTACTGTTATTGGGACTATTTA 100
G G T F R G N V R G T P I C L F F . I H S N M Y P L M R Q . P . . M
GTTCAATAATATTGAAAAAGGAAGATGATGATTCACATTTCCGCTGCGCCCTTATCCCTTTTTCGGCAATTTGCCCTTCTGTTTTCCTCAC
CSAAGTTATTATAACTTTTCTCTCACTACTCATAAGTTGTAAGGCACACGCGGAATAAGGAAAAACCGCGTAAACGGAAGGACAAAAACGAGTG 200
L Q . Y . K R K S M S I Q H F R V A L I P F F A A F C L P V F A H
CDAGAAACGCTGGTGAAGTAAAGATGCTGAAGATCAGTTGGGTGCAGAGTGGGTACATCGAATGGATCTCAACAGCGGTAAAGATCCTTGAGAGTT
GGTCTTTGGGACCACTTTCATTTCTACGACTCTAGTCAACCCACGTCCTCACCAGTATGATGACCTAGAGTTGTCGCCATTTAGGAACCTCTCAA 300
P E T L V X V K D A E D O L G A R V G Y I E L D L N S G K I L E S
TTGCCCCGAAGAAGCTTTTCCAATGATGAGCACTTTTAAAGTTCTGCTATGTCGGCGGCTATTATCCGCTATTGACGCGGGCAAGAGCAACTCGGTCG
AAGCGGGCTCTTGCAAAAGGTTACTACTCGTGAAATTTCAAGACGATACACCGCGCCATAATAGGGCATAACTGCGGCCCTTCTCGTTGAGCCAGC 400
F R P E E R F P M H S T F K V L L C G A V L S R I D A G Q E Q L G R
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R I H Y S Q N O L V E Y S P V T E K H L T D G H T V R E L C S A A
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I T H S O N T A A N L L L T T I G G P K E L T A F L H N M G D H V
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T R L D R V E P E L N E A I P N D E R D T T N P V A M A T T L R K L
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A G V F I A D K S G A G E R G S R G I I A A L G P O G K P S R I V
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V I Y T T G S O A T M D E R N R O I A E I G A S L I K H V . L S D O
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V Y S Y I L . I D L K L H F F K R I . V K I L F D N L M T K I P
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fig 5-3 pLM1 (1 > 8285) Site and Sequence

Page 2

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A S G D K S C L T G L D S R R . L P D K A Q R S G . T G G S C T Q

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P S L E R T T Y T E L R Y L O R E L . E S A T L P E G R K A D R Y P

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E R R F L . C S S G G R S L V K N A S N A A F L R F L A F C V P F

CTCACATGTTCTTCTTCGCTTATCCCTGATTCTGTGGATAACCGTATTACCGCTTTGAGTGAGCTGATACCGCTCGCGCAGCCGAACGACCGAGCG
GAGTGACAAAGGAGCGCAATAGGGGAC TAAGACACCTATTGGCATAATGGCGGAAAC TCACCTGACTATGGCAGCGCGCTCGGCTTGTGCTGCTCG
A H M F F P A L S P O S V D N R I T A F E . A D T A R R S R T T E R

CAUCGAGTCAGTGAGCGAGGAAGCGGAAGAGCGCCCAATACGCAACCGCTCTCTCCCGCGCTTGGCCGATTTCATTAATGCAGCTGGCAGCAGAGTTT
GTGCTCTAGTCACTCGCTCTTTCGCTTCTCGCGGTATGCGTTTTCGCGGAGAGGGGCGCGCAACCGCTTAAGTAATACGTCGACCTGCTGCTCAAA
G S E S V S E E A E E R P I R K P P L P A R V P I H . C S V H D R F

CCGACCTGGAAGCGGCGAGTGAGCGCAACGCAATTAATGTAGTGTAGCTACCTCATATTAGCCACCCAGGCTTTACACTTTATGTCTTCGCGCTGTATGT
GGGCTGACCTTTTCGCGCTCTACCTCGCTTTCGCTTAATTACACTCAATCGAGTGAGTAATCCGTTGGGTTCGAAATGTGAAATACGAAGCGCGAGCATA
P D V K A G S E R N A I N V S . L T H . A P Q A L H F H L P A R M

TGTGTGGAATGTGAGCGGATAACAATTTACACAGGAAACAGCTATGACCATGATTACGCCAAGCGCGCAATTAACCTCTACTAAAGGGAACAAAGCT
ACACACCTTAACACTCGCTATGTTAAAGTGTGCTTTTCTGSA*ACTGGTACTAATGCGGTTCGCGGTTAATTGGGAGTGATTCCCTTGTGTTTCGA
L C G I V S G . Q F H T G N S Y D H O Y A K R A I N P H . R E Q K L

GGGTACCGGGCCCCCTCGAGGTGAGCTATCGATAGCTTGTATGCAATTCCTGCAGCCCTGCTCTTCAGCCAGATGCTGGACCCAGAGTCCAG
CCATGCGCGCGGGGGAGCTCCAGCTGCTATAGCTATTCGAACTA*AGCTTAAGGAGCTTCGGGGAGCAGAAGTCTGCTACGACCTGGGTCTCAGGGTC

insert pLM1

ORF pLM1

G T G P P L E V D G I D K L D ! E F L O P L L F S Q N L O P E S Q

AGAAAGAGGACAGTGCAGAAATGCTCGATCTCGGCGAGAACTGGAAGAGACCATGTCCAGCTCGGAGGGTCCAGGCTGACACAGCTCCCTGGAGA
CTTTCTCTCTGCTACGCTTACAGGACCTAGAGGGCTCTTGGACTCTCTGGTACAGGTGCGAGCTCCAGGGTCCACTGAGTGTGAGGGGACCTCT

insert pLM1

ORF pLM1

R K R T V Q N V L D L R Q N L E E T M S S L R G S Q Y T H S S L E

TACCTGCTACGACAGCATGATGCCAAGCCAGCGAGCTGTTCAGCTCTCCAAACCGCTGCTCCCTCTGTCTATGCGCTATGCGCAGTCCAGTCCGG
ACTGAGCAGATGCTGTGCTACTACGGTTGCGTCTCGCTGCTACAGCTTGGAGGTTGGCGAGCAGGGGAGACAGTACCGCATACCGTCAAGTCAAGCGC

insert pLM1

ORF pLM1

M T C Y D G D O A N P R S V S S L S N R S S P L S V R Y G Q S S P R

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Fig 54 pLM1 (1 > 8285) Site and Sequence

GC TGCAGGCTGGTGCACGCGCCCTCTGTGGGTGGGAGCTTCCCTCTGAGGGGACGCCCGCCTGTACATGCACGGCGAACGGGCCCAC TACTCCACACC 2600
CGACGTCCGACCACTGCGCGGGAGACACCCACCCCTCGACGGGAGCCCTCCCTTCCGGGGGACCATGTACGTGCCGCTTGCCCGGGTGATGAGGGTGTGG
-----insert pLM1-----
-----ORF pLM1-----
L Q A G D A P S V G G S C R S E G T P A V Y M H G E R A H Y S H T
ATGCCCATGCGCAGCCCCAGCAAGCTCAGCCATATCTCCCTCTGAGGCTGGTGAATCCCTGGACTCGGATGAGGTGGACC TCAAGTCCGGCTACATGA 2700
TACGGGTACGCGTCGGGGTCGTTCGAGTCGGTA TAGAGGGGCGACCTCGACCAGCTTAGGGACCTGAGCCTACTCCACCTGGAGTTCAGGCCGATGACT
-----insert pLM1-----
-----ORF pLM1-----
M P M R S P S K L S H I S S L E L V E S L O S D E V D L K S G Y M
GCGACAGTGACCTCATGGGCAAGACCATGACGGAGGATGATGATCATCATACACCGGCTGGGATGAAAGCAGCTCCATCAGTAGTGGACTCAGCGATGCCCTC 2800
CGCTGTCACTGGAGTACCCGTTCTGGTACTGCCCTCCTACTCTGTAGTGATGGCCGACCTACTTTCTGTCGAGGTAGTCACCTGAGTCGCTACGGAG
-----insert pLM1-----
-----ORF pLM1-----
S D S D L M G K Y T M T E O C D I T T G V D E S S S I S S G L S D A S
AGACAATCTCAGTTTCAAGAAGTCAATGCCAGCTCCTCACTCAACTCCCTCCCAAGTACTCCCATGCTTCTCGCAGGAAC TCAACAATAGTGCTACGC 2900
TCTGTTAGAGTCAAGTCTTCTTAAGTACGGTCGAGGAGTGAATGAGGGAGGGTTCATGAGGGTGACGAAGAGCGCTCTTGAGTTGTTATCACGATGCG
-----insert pLM1-----
-----ORF pLM1-----
D N L S S E E F N A S S S L N S L P S T P T A S R R N S T I V L R
ACAGACTCAGAGAAGCGCTCACTGGCAGAAAGTGGGCTGAGTGGTTAGTGAATCAGAGGAGAAAGCCCTAAAAAAGTGGAGTACGACAGTGGTAGCC 3000
TGTC TGAGTCTCTTCGCGAGTGACCGTCTTTCACCCGACTGACCAATCACTTAGTCTCTCTTTGCGGGATTTTTGACCTCATGCTGTCACCATCGG
-----insert pLM1-----
-----ORF pLM1-----
T D S E K R S L A E S G L E V F S E S E E K A P K K L E Y D S G S
TGAAGATGGAACCTGGGACTTCTAAGTGGCAGAGGAGTGGGCTGAGTGGTGAATGATTCATCCAAGGGTGGGAGAC TGAAGAGCCCATAGCCTGGG 3100
ACTCTACCTTGGACCTTGAAGATTCACCGCCTCCCTGCGGAGTCTGACACTACTAAGTAGGTTCACCTCTTGAC TTTTTCGGGTAGTCGGACCC
-----insert pLM1-----
-----ORF pLM1-----
L K M E P G T S K V R R E P E S C D D S S K G G E L K K P I S L G
GCAACCTGGTTCCTGAGAAGGGCAAGACCCACCTGCTCTTGAAGTTCCTCCATCACTCACACAGCCAGAGTGCCT TCAAGTCGCGAGGCAACCT 3200
GCTGGGACCAAGGACTTCTTCCCGTCTCTGGGTGGATACCTGAGTGAAGGGGGTAGTGAGTGTGTCGGGTCTCACGGGAGTTTCAGGCTCCGTTTGGA
-----insert pLM1-----
-----ORF pLM1-----
H P G S L K K G K T P P V L V T S P I F H T A Q S A L K V A G K P
GAGGCAAGCTACAGACAAGGTAAGCTGCAAGTGAAGTACTGAGTCAAGCTCTCTCTGATGCTGGTGGGACCGCTGAGTGA TGC TAAGA 3300
GCTGCTTTCGATGCTGTTTCCATTCGAACCTCACTTCTTATGAGTGAAGTTCGGAGGAGGAGACTACGACCAAGCCCTGGCTGAC TCACTACGATCT
-----insert pLM1-----
-----ORF pLM1-----
E I K A F D K G K L A V K K I L Q R S S S D A G R C P L S D A K

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[illegible]

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[illegible]

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fig 54 pLM1 (1 > 8285) Site and Sequence

Page 1

ATGAGGGACAGGTTCTTGGTGTGTACCTTTGAGAACTTCCTAGGAAGGAATGGTGGGGTGGCGTTTGGGAACCTTGCCCCCTAAACACATTACTGGC
TCTCCCTGTCCAAGAACACGACATGGAAACTCTTGAAGGATCTTCCCTTACCACCCACCGCAACCCCTTGAACACGGGGATTGTGTAATGACCG 7400

insert pLM1

G G T G S V C C T F E N F L G R N G G V A F G N L C P L N T F T G

CTCTCTAATGACTTTGGGAAAAGATGATTCTGGGTCTTCCCTTGACTTCTGTTTCAATTACAACTCCTGGGCTTCTGGGGAGGGGTTCAAGAAA
GAGGAGATTACTGAAACCCCTTTCTACTAAGACCCATAAGGGAAGTGAAGAACAAAGTTAATGTTTGGAGCCCGAAAGACCCCTCCCAAGTCTTTT 7500

insert pLM1

L L . . L V G K D D S G S F P . L L V S I T N S V A F V G G V Q K

CATCAAACTGACGAGTTCCTCCGAATTCAGCTTTCACCTAACAGGCTGAACCTTGCTCAAAAGAAGCCGAATTCAGCACACTGGCGCCGTTACT
GTAGTTTGTGACGTCGTCAAGGGGCTTAAGTCGAACCTGAATTGGTCCGACTTGAACGAGTTTCTTCGGCTTAAGGTCGTGTGACCCCGGCAATGA 7600

insert pLM1

T S K H C S S S P E F S L D L Y R L N L L K R S R I P A H V R P L L

AGTTCTAGAGCGGCGCCACCGCGGTGGAGCTCCAATTGCGCTATAGTGAGTCGTATTACGCGCGCTCACTGGCGGTGTTTTACAACTGTCGTACTGG
TCAAGATCTCGCGCGGTGGCGCCACCTCGAGGTTAAGCGGATATCACTCAGCATAATGCGCGGAGTGACCGGCAGCAAAATGTTGACGCACTGACC 7700

3

V L E R P P P R V S S N S P Y S E S Y Y A R S L A V V L O R R D V

GAAAACCTTGGCGTTACCAACTTAATCGCCTTGACGACATCCCTTTTCGCCAGCTGGCGTAATAGCGAAGAGGCGCCGACCGATCGCCCTTCCCAAC
CTTTGGGACCGCAATGGTGTGAATFAGCGGAACGCTCTAGTGGGGAAGCGGTCGACCGCATTATCGCTTCTCGGGCGTGGCTAGCGGGAAGGGTTG 7800

E N P G V T Q L N R L A A H P P F A S V R N S E E A R T D R P S Q

AGTTGCGCAGCTGAATGGCGAATGGGACGCGCCCTGTAGCAGCGCATTAGCGCGCGGGGTGTGGTGGTACGCGCAGCTGACCGCTACACTTGCCAG
TCAACGCGTCGGACTTACCGCTTACCTTGCGCGGGACATCGCCCGTAATTCGCGCGCCACACCAATGCGCTCGCACTGGCGATGTGAACGGTC
Q L R S L N G E V D A P C S G A L S A A G V V V T R S V T A T L A S 7900

CGCCCTAGCGCCGCTCTCTTCGCTTCTTCCCTTCTCTTCTGCGCAGCTTCGCGGCTTCCCGTCAAGCTCTAAATCGGGGCTCCCTTTAGGGTTC
CTGGATCGCGGCGAGGAAACGAAAGAGAGGAAAGGAAAGAGCGGTGCAAGCGGCGGAAAGGGGAGTTTCGAGATTAGCCCCGAGGGAATCCCAAG
A L A P A P F A F F P S F L A T F A G F P R O A L N R G L P L G F 8000

CTATTTAGTGCTTACGGCACCTCGACCCCAAAAACCTTATTAAGTGGTTCACGTAGTGGGCCATCGCCCTGATAGACGTTTTTCGCCCTTTGA
CTTAATCAGGAAATGCGGTGGAGCTGGGGTTTTTGAATTAATCCCACTACCAAGTGCATCACCCTGAGCGGACTATCTGCCAAAAGCGGGAATCT
R F S A L R H L D P K K L D . G D G S R S G P S P . . T V F R P L 8100

CTTTGAGTCCACGTTCTTTAATAGTGGACTCTTGTCTCAAACTGGAACAACTCAACCCCTATCTCGGTCTATTCTTTTGATTATAAGGGATTTTGGC
GCAACCTCAGGTGCAAGAAATFATCAGCTGAGAACAAGTTTGAACCTTGTGTGAGTTGGGATAGAGCCAGATAAGAAAATAAATATTCCTTAAACGG 8200

I L E S T F F N S G L L F D T G T T L N P I S V Y S F O L . G I L P

CTTTTCGGCTATTGGTTAAAAAATGAGCTGATTTAACTAAAAATTAACGCGAATTTTAAACAAATATTAACGCTTACAATTTAG
CTAAAGCGGATAACCAATTTTCTACTGCACTAAATTTCTTAAATTCGCTTAAATTTGTTTATAATTCGCAATGTTAAATC 3295

: S A Y V L K H E L I . : < F N A N F N K I L T L T I .

SEQUENCE ID PCT

Page 1

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fig 34 pLM4 (1 > 10070) Site and Sequence

Enzymes : 100 of 146 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

100
TAGTTATTAATAGTAATCAATTACGGGGTCATTAGTTTCATAGCCCATATATGGAGTTCCCGGTTACATAAATTACGGTAAATGGCCCGCTGGCTGACCC
ATCAATAATTATCATTAGTTAATGCCCCAGTAATCAAGTATCGGGTATATACCTCAAGGCGCAATGTATTGAATGCCATTTACCGGGCGGACCGACTGGC
PCMV
L L I V I N Y G V I S S . P I Y G V P R Y I T Y G K V P A V L T
200
CCCCACGACCCCGCCCATTCAGCGTCAATAATGACGTATGTTCCCATAGTAACGCCAATAGGGACTTTCATTGACGTCATGGGTGGAGTATTTACGGT
GGGTTGCTGGGGCGGGTAACGTCAGTTATTACTGCATACAAGGGTATCATTGCGGTTATCCCTGAAAGGTAACGTCAGTTACCCACCTCATAAATGCCA
PCMV
A Q R P P P I D V N N D V C S H S N A N R D F P L T S M G V F T V
300
AAACTGCCCACTTGGCAGTACATCAAGTGTATCATATGCCAAGTACGCCCCCTATTGACGTCATGACGGTAAATGGCCCGCTGGCATTATGCCAGTA
TTTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTCATGCGGGGATAACTGCAGTTACTGCCATTTACCGGGCGGACCGTAATACGGGTCAT
PCMV
N C P L G S T S S V S Y A K Y A P Y . R Q . R . M A R L A L C P V
400
CATGACCTTATGGGACTTTCCTACTTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGATGCGGTTTGGCAGTACATCAATGGGCGTGGA
GTACTGGAATACCTGAAAGGATGAACCGTCATGTAGATGCATAATCAGTAGCGATAATGGTACCACACGCGCAAAACCGTCATGTAGTTACCCGACCT
PCMV
H D L M G L S Y L A V H L R I S H R Y Y H G D A V L A V H Q V A V
500
TAGCGGTTTGACTCAGGGGATTTCCAAGTCTCCACCCCATTCAGCGTCAATGGGAGTTTGTTTTGGCACCAAAATCAACGGGACTTTCACAAATGTCGTA
ATCGCAAACTGAGTGCCCTAAAGGTTCAAGGTTGSGGTAACGTCAGTTACCTCAAAACAAACCGTGGTTTGTAGTTGCCCTGAAAGGTTTACAGCAT
PCMV
I A V . L T G I S K S P P H . R Q V E F V L A P K S T G L S K M S .
600
ACAACTCGCCCCCATTCAGCAAAATGGGCGGTAGGCGTGTACGGTGGGAGGTCTATATAAGCAGAGCTGGTTTAGTGAACCGTCAGATCCGCTAGCGCTA
TGTGAGGCGGGTAACGCGTTTACCCGCCATCCGACATGCCACCTCCAGATATATTCGTCTCGACCAAACTCACTTGGCAGTCTAGGCGATCGCGAT
PCMV
Q L R P I D A N G R . A C T V G G L Y K Q S V F S E P S D P L A L
700
CCGGTCGCCACCATGGTGAGCAAGGGCGAGGAGCTGTTACCGGGGTGGTGCCCATCTGGTTCGAGCTGGACGGCGACGTAACGGCCACAAGTTCAGCG
GGCAGCGGTGGTACCACTCGTTCCCGCTCCCTCGAAGTGGCCCCACCGGGTAGGACCAAGCTCGACCTGCCGCTGCATTTGCCGGTGTTCAGTCCG
EGFP
P V A T M V S K G E E L F T G V V P I L V E L D G D V N G H K F S
800
TGTCCGGCGAGGGCGAGGGCGATGCCACCTACGGCAAGCTGACCTGAAGTTGATCTGCACACCGGCAAGCTGCCCGTGCCCTGGCCCACTCTGTGAC
ACAGGCCGCTCCCGCTCCCGCTACGGTGGATGCGCTGACACGCGGACCTGAGGACTTCAAGTAGACGTGGTGGCCGTTCGACGGGCACGGGACCGGTGGGAGCACTG
EGFP
V S G E G E G D A T Y G K L T L K F I C T T G K L P V P V P T L V T
900
CAGCGTGACCTACGGCGTGCAGTGCCTCAGCGCTACCGGACCATGAAGCAGCAGCACTCTTCAAGTCCGCCATGCCCGAAGGCTACGTCAGGAG
GTGGGACTGGATGCCGACCTCAGCAAGTGGGCAAGGCGCTGGTGTACTTGGTGGTCTGAAGAAGTTCAGGCGGTACGGGCTTCGATGCAAGTCTCTC
EGFP
L T T G V G C F S R . P D H N K O H D F F K S A M P E G Y V Q E

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fig 34 pLM4 (1 > 10070) Site and Sequence

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CGCACCATCTTCTTCAAGGACGACGGCAACTACAAGACCCGCGCCGAGGTGAAGTTCGAGGGCGACACCCCTGGTGAACCGCATCGAGCTGAAGGGCATCG
CGGTGGTAGAAGAAGTTCCTGCTGCCGTTGATGTTCTGGGCGCGGCTCCACTTCAAGCTCCCGCTGTGGGACCCTTGGCGTAGCTCGACTTCCCGTAGC
EGFP
RTIFFKODGN YKTRA EVKFEGDTLVHRIELKGI
ACTTCAAGGAGGACGGCAACATCTGGGGCACAAGCTGGAGTACAAC TACAACGCCACAACGCTATATCATGGCCGACAAGCAGAAGAACGGCATCAA
TGAAGTTCTCTGCGGTTGTAGGACCCCGTGTTCGACCTCATGTTGATGTTGTCGGTGTTCAGATATAGTACCAGCTGTTCGTCTCTTGGCGTAGTT
EGFP
DFKEDGNILGHKLEYN YNSHN VYIMADKQKNGIA
GGTGAATTCAGATCCGCCACAACATCGAGGACGGCAGCGTGCAGCTCGCCGACCACTACCAGCAGAACACCCCATCGGCAGCGGCCCGTGTCTGT
CCACTTGAAGTTCTAGCGGTTGTAGCTCCTGCCGTGCGACGTCGAGCGGCTGGTGTGGTCTGTCTTGTGGGGTAGCCGCTGCCGGGGCACGACGAC
EGFP
VNFKIRHNIE DGSVQLADHYQQNTPTIGDGPVLL
CCCGACAACCACTACCTGAGCACCCAGTCCGCCCTGAGCAAGACCCCAACGAGAAGCGCATCACATGGTCTGTCTGGAGTTCGTGACCGCGCCGGGA
GGGCTGTGGTGTGGACTCGTGGGTCAGGCGGGACTCGTTTCTGGGGTGTCTCTTCGCGCTAGTGTACCAGGACGACCTCAAGCACTGGCGGGCGCCCT
EGFP
PDNH YLS TQS ALSKDPNEKR D H M V L L E F V T A A G
TCACTCTCGGCATGGACGAGCTGTACAAGTCCGACTCAGATCTCGAGCTCAAGCTTCGAATCTGCAGTCGATAAGCTTGATATCGAATTCCTGCAGCC
AGTGAGAGCCGTACCTGCTCGACATGTTTCAGGCTTGAGTCTAGAGCTCGAGTTCGAAGCTTAAGACGTCAGCTATTTCGAATATAGCTTAAGGACGTCGG
EGFP
ITLGMDELYKSSGLRSRAQASNSAVDKLDIEFLQP
CCTGCTCTTCAGCCAGATGCTGGACCCAGAGTCCCAAGAGAGGACAGTGCAGAAATGCTCTGGATCTCCGGCAGAACCTGGAAGAGACCATGTCCAGC
GGACGAGAAGTCGGTCTACGACCTGGGTCTCAGGGTCTCTTCTCTCTGTCACGCTTACAGGACCTAGAGGCGCTCTGGACCTCTCTGTTACAGGTCT
insert pLM1
ORF pLM1
LLFSQMLDPE SQRKRTVQNVLDLRLONLEETMS S
CTGCGAGGGTCCCAGGTGACTCACAGCTCCCTGGAGATGACCTACGACAGCGATGATGCCAACCCACGACGCTGTCCAGCTCTCCAACCGCTCG
GACGCTCCAGGGTCCACTGAGTGTGAGGGACCTCTATGAGCATGCTGTGCTTACGGTGGGTGCGTCCGACAGGTGCGAGAGGTTGGCGAGCA
insert pLM1
ORF pLM1
LRGSQVTHSSLEMTCYDSDDANPRSVSSLSNRS
CCCCCTGTGTCATGGCGCTATGGCCAGTCCAGTCCGCGCTGACGGCTGGTGACGCGCCCTCTGTGGGTGGGAGCTGCCGCTCGGAGGGGACGCCCGCT
GGGAGACAGTACCGCATACCGGTACGGTCAAGGCTGAGGCTGACGCGGAGACACCCACCTTCGACGCGAGGCTCCCCCTGCGGGCGGAC
insert pLM1
ORF pLM1
PLSRIRISSSRLDAGDAPSVGGSCRSSEGTDAV

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fig 34 pLM4 (1 > 10070) Site and Sequence

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G T A C A T G C A C G G C G A A C G G G C C C A C T A C T C C C A C A C C A T G C C C A T G C G C A G C C C C A G C T C A G C C A T A T C T C C C G C C T G G A G C T G G T C G A A T C C C T G
C A T G T A C G T G C C G C T T G C C C G G G T G A T G A G G G T G T G G T A C G G T A C G C G T C G G G T C G T T C G A G T C G G T A T A G A G G G C G G A C C T C G A C C A G C T T A G G G A C

insert pLM1

ORF pLM1

Y M H G E R A H Y S H T M P M R S P S K L S H I S R L E L V E S L

G A C T C G G A T G A G G T G G A C C T C A A G T C C G G C T A C A T G A G C G A C A G T G A C C T C A T G G G C A A G A C C A T G A C G G A G G A T G A T G A C A T C A C T A C C G G C T G G S A T G
C T S A G C C T A C T C C A C C T G G A G T T C A G G C C G A T G T A C T C G C T G T C A C T G G A G T A C C C G T T C T G G T A C T G C C T C C T A C T A C T G T A G T G A T G G C C G A C C C T A C

insert pLM1

ORF pLM1

D S D E V D L K S G Y M S D S D L M G K T M T E D D D I T T G V D

A A G C A G C T C C A T C A G T A G T G G A C T C A G C G A T G C C T C A G A C A A T C T C A G T T C A G A A G A A T C A A T G C C A G C T C C T C A C T C A A C T C C C T C C C A A G T A C T C C
T T T C G T C G A G G T A G T C A T C A C C T G A G T C G C T A C G G A G T C T G T T A G A G T C A A G T C T T C T T A A G T T A C G G T C G A G G A G T G A G T T G A G G G A G G G T T C A T G A G G

insert pLM1

ORF pLM1

E S S S I S S G L S D A S D N L S S E E F N A S S S L N S L P S T P

C A T G C T T C T C G C A G G A A C T C A A C A A T A G T G C T A C G C A C A G A C T C A G A G A A G C G C T C A C T G G C A G A A A G T G G G C T G A G C T G G T T T A G T G A A T C A S A G G A G
G T S A C G A A G A G C G T C C T T G A G T T G T T A T C A C G A T G C S T G T C T S A G T C T C T C G C G A G T G A C C G T C T T C A C C C G A C T C G A C C A A A T C A C T T A G T C T C C T C

insert pLM1

ORF pLM1

T A S R R N S T I V L R T D S E K R S L A E S G L S W F S E S E E

A A G C C C C T A A A A A C T G G A G T A C G A C A G T G G T A C C C T S A A G A T G G A A C C T S G G A C T T C T A A G T G G C G G A G G G A G C G G C C T G A S A G C T G T S A T S A T T C A T
T T T C G G G A T T T T T T G A C C T C A T G C T G T C A C C A T C G S A C T T C T A C C T T G G A C C C T G A A G A T T C A C C G C C T C C C T C G C C G G A C T C T C G A C A C T A C T A A G T A

insert pLM1

ORF pLM1

K A P K K L E Y C S G S L K M E P G T S K V R R E R P E S C D D S

G C A A G G G T G G A G A A C T G A A A A G C C C A T C A G C C T G G S C C A C C C T G G T T C C C T G A A G A A G S C A A G A C C C C A C C T G T G G C T G T A A C T T C C C C A T C A C T C A
G G T T C C C A C C T C T T G A C T T T T T C G G S T A S T C G G A C C C S S T G G S A C C A A G G S A C T T C T C C C S T T C T G G G G T G G A C A C C G A C A T T G A A G G G S S T A S T G A G T

insert pLM1

ORF pLM1

G G E L K K P I S L S - P G S L K K G K T P P V A V T S P I T H

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fig 34 pLM4 (1 > 10070) Site and Sequence

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CACAGCCCAGAGTGGCC TCAAAGTCGCASGCCAAACCTGAGGGCAAAGCTACAGACAAGGGTAAGCTTGCAGTGAGGAATACTGGGGTCCAAACGCTCCTCC
GTGTGGGGTCACAGGGAGTTTACAGCGTCCGTTTGGACTCCCGTTTCGATGTCTGTITCCCATTCGAACGTCACITTCATGACCCGAGGTTGCSAGGAGG

2400

insert pLM1

ORF pLM1

T A O S A L K V A G K P E G K A T D K G K L A V K N T G L O R S S

TCTGATGCTGGTCGGGACCGCTGAGTGATGCTAAGAAGCCCCCTCGGGCATTGCTCGCCCCCTCCACTTCGGGATCCTTCGGCTACAAGAAGCCTCCTC
AGACTACGACCAGCCCTGGCGGACTCAC TACGATTCTTCGGGGGAGCCCGTAACGAGCGGGGAGGTGAAGCCCTAGGAAGCCGATGTTCTTCGGAGGAG

2500

insert pLM1

ORF pLM1

S D A G R D R L S D A K K P P S G I A R P S T S G S F G Y K K P P

CTGCCACAGGCACAGCCACTGTCTATGCAAAC TGGTGGTTACAGCCACTCTCAGCAAGATCCAGAAGTCTCAGGCATCCCTGTCAAGCCAGTAAATGGGCG
GACGGGTGTCGGTGTGGTGACAGTACGTTTGACCACCAAGTCGGTGAGAGTCGTTCTAGGTCTTCAGGAGTCCGTAGGGACAGTTTCGGTCAATTACCCGC

2600

insert pLM1

ORF pLM1

P A T G T A T V M O T G G S A T L S K I O K S S G I P V K P V N G R

CAAGACTAGCTTAGATGTTTCCAACAGCGCAGAGCCAGGATTCCTGGCTCCTGGAGCCCGTTCTAACATCCAGTACCGCAGCCTGCCCCGGCCAGCCAAAG
GTCTGATCGAATCTACAAAGGTTGTCGGTCTCGGTCC TAAGGACCGAGGACCTCGGGCAAGATTGTAGGTCATGGCGTCGGACGGGCGGGTTCGTTTC

2700

insert pLM1

ORF pLM1

K T S L D V S N S A E P G F L A P G A R S N I O Y R S L P R P A I

TCAAGTTCTATGAGCGTGACCGGCGGGGGGGTGGACCTCGCCCTGTGAGCAGCAGCATTGACCCCAAGTCTCTCAGCACCAAGCAGGGAAGGCTTACGG
AGTTCAAGATACTCGCACTGGCCGCGCCGCCCACTGGAGCGGGGACACTCGTCGTCGTAAC TGGGGTCAGAGGAGTCGTGGTTCGTCCTCCGGGAATGCG

2800

insert pLM1

ORF pLM1

S S S M S V T G G R G G P R P V S S S I D P S L L S T K O G G L T

CTTCCAGACTGAAGGAGCCTACCAAGGTAGCCAGTGGGCGGACCCTCCAGCCCCCTGTCAATCAGACAGATCGGGAAAAGGAGAAGGCCAAAGCCAAGG
GAAGGTCGACTTCTTCGGATGTTTCCATCGGTCACCCGCTGGTGAGGTCGGGGACAGTTAGTCTGTC TAGCCCTTTTCTCTTCGGGTTTCGGTTCCG

2900

insert pLM1

ORF pLM1

F S R L K E P T K V A S G R T T P A P V H O T C R E X E K A A A I

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AGTGGCC TTGGACTCAGACAACATC TCCTTGAAGAGTATTGGCTCCCCAGAGAGTAC TCCCAAGAACCAAGCAAGCCACCCCAAGCCACCAAGCTGGCA
TCACCGGAACCTGAGTCTGTTGTAGAGGAACCTCTCATAACCGAGGGGTCCTCATGAGGGTTCCTTGGTTCGTTCGGTGGGGTCTCGGTGGTTCGACCGT 300

insert pLM1

ORF pLM1

V A L D S D N I S L K S I G S P E S T P K N Q A S H P T A T K L A

GAGCTGCCACCAACCCCTCTCAGGGCCACAGCGAAGAGCTTTGTCAAACCAACCTCACTAGCCAATCTTGACAAGGTCAACTCCAACAGTCTGGATCTAC
CTCGACGGTGGTTGGGGAGAGTCCCGGTGTCGCTTCGAAACAGTTTGGTGGGAGTGATCGGTTAGAACTGTTCAGTTGAGGTGTGCAGACCTAGATG 310

insert pLM1

ORF pLM1

E L P P T P L R A T A K S F V K P P S L A N L D K V N S N S L D L

CATCATCCAGTGATACCACCATGCTTCAAAGGTCCCAGATCTGCATGCTACAAGCTCAGCATCTGGGGGCCCTCTCCCTTCCTGCTTCACCCCAAGTCC
GTAGTAGGTCACTATGGTGGGTACGAAGTTTCCAGGGTCTAGACGTACGATGTTTCGAGTCGTAGACCCCGGGAGAGGGAAGGACGAAGTGGGGTTCAG 320

insert pLM1

ORF pLM1

P S S S D T T H A S K V P D L H A T S S A S G G P L P S C F T P S P

GGCACCCATCCTCAATATTAACCTCAGCCAGCTTCTCCAGGGCCCTGGAGCTAATGAGTGGTTTCAGTGTGCCAAAAGAGACCCGCATGTACCCCAAACTC
CCGTGGGTAGGAGTTATAATTGAGTCGGTCGAAGAGGGTCCCGGACCTCGATTACTACCAAAAGTCACACGGTTTTCTCTGGGCGTACATGGGGTTTGA 330

insert pLM1

ORF pLM1

A P I L N I N S A S F S Q G L E L M S G F S V P K E T R M Y P K L

TCAGGCCCTGCACAGGAGCATGGAGTCCCTCCAGATGCCAATGAGCTCCCAAGTGCCTTCCCAAGCAGTACTCCCGTCCCAACCCACCTGCTCCCGCTG
AGTCCGGACGTGTCCTCTACCTCAGGGAGGTCTACGGTTACTCGGAGGGGTACGGAAGGGGTCTGTCATGAGGGCAGGGGTGGGGTGGACGAGGGGGAC 340

insert pLM1

ORF pLM1

S G L H R S M E S L Q M P M S L P S A F P S S T P V P T P P A P P

CTGCTCCCAAGAGAAGAGAGCGGAAGAGCTGACTTGGAGTGGAAAGCCCAAGAGCTGGGCAACTGGACAGTAAACAGCGGGATCGGAACACTCTTCGCAA
GAGGAGGGGTGCTCTCTCTGCTCTCTCGACTCAACCTCACCTTCGGGGTCTCGACCCGTTGACCTGTGATTAGTCGCGCTAGCCTTGTGAGAAGGGTT 350

insert pLM1

ORF pLM1

A A P T E E E T E E L T V S G S P R A G Q L D S N Q R Q R N I L F I

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GAAAGGGCTCAGGTACCAGCTTCAGTCCCAGGAGGAGACCAAGGAGAGGCGACATCCCATACCATGGTGGGCTGCCGTAATCCGATGACCAGTCAGAG
CTTTCCCGAGTCCATGGTCAAGTCAGGGTCTCTCTGGTTCTCTCCGCTGTAAGGGTATGGTAACCAACCCGACGGACTTAGGCTACTGGTCAGTCTC 360

insert pLM1

ORF pLM1

K G L R Y Q L O S Q E E T K E R R H S H T I G G L P E S D O Q S E

CTGCCCTTCTCCCCCTGCACCTCCCATGTCTCTGAGTGCAAGGGCCAACCTACCAACATAGTGAGTCCCCTGCGGCCACCCAGCCAAGAATCACCCGCT
GACGGAAAGGGGGACGTGAAGGGTACAGAGACTCACGTTTCCCGGTTGAATGGTGTATCACTCAGGGTGACGCCGGTGGTGGGTTCTTAGTGGGGCA 370

insert pLM1

ORF pLM1

L P S P P A L P M S L S A K G Q L T N I V S P T A A T T P R I T R

CCAACAGCATCCCCACCCACGAGGCGGCCCTTCGAGCTGTACAGCGGCTCCCAATGGGGAGCACCTGTCCCTGGCCGAGAGACCCAAGGGAATGATTCC
GGTTGTCTGAGGGGTGGGTGCTCCGCCGAAGCTCGACATGTGCCGAGGGTTTACCCCTCGTGGGACAGGGACCGGCTCTCTGGGTTCCCTTACTAAGC 380

insert pLM1

ORF pLM1

S N S I P T H E A A F E L Y S G S Q M G S T L S L A E R P K G M I R

GTCAGGATCCTTCCGAGACCCACGGACGATGTTACGGCTCAGTGCTGTCCCTGGCCCTCCAGTGCCCTCCTCCACCTACTCCTCAGCTGAGGAGAGGATG
CAGTCTTAGGAAGGCTCTGGGGTGCCTGCTACAAGTGCCGAGTCAACGACAGGGACCGGAGGTCACGGAGGAGGTGGATGAGGAGTCGACTCCTCTCCTAC 390

insert pLM1

ORF pLM1

S G S F R D P T D D V H G S V L S L A S S A S S T Y S S A E E R M

CAATCTGAGCAAAATCCGGAAGCTTCGTAGGGAACGGGAATCATCCAGGAAAAAGTGGCCACCTTGACGCTCAGCTTCTTGCCAATGCTAATCTGGTGG
GTTAGACTGGTTTAGGCTTCGAAGCATCCCTTGACCTTAGTAGGGTCTTTTTCACCGGTGGAACGTCAGAGTCGAAAGACGGTTACGATTAGACCAC 400

insert pLM1

ORF pLM1

D S E Q I R K L R R E L E S S O E K V A T L T S Q L S A N A N L V

CTCCTTTTGAGCAGAGCCTGGTGAATATGACATCCCGCTGCGACACCTGGCAGAGACGGCCGAGGAGAAGGACACTGAGCTGCTGGATTTGCGAGAAAC
GACGAAAACCTCGTCTCGGACCACTTATACTGTAGGGCGSACGCTGTGGACCGTCTCTGCCGGCTCCTCTCTCTGTGACTCGACGACCTAAACGCTCTTG 410

insert pLM1

ORF pLM1

A A F E Q S L V N M T S R L R H L A E T A E E K O T E L L O L R E I

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CATAGACTTTCTGAAGAAAAGAACTCTGAGGCCCAAGCAGTCATTACGGGAGCCCTTAATGCCTCAGAAACCACACCCAAAGAACTTCGGATCAAGAGA
GATCTGAAAGACTTCTTTTCTTGAGACTCCGGGTCCTCAGTAAGTCCCTCGGGAATTACGGAGTCTTTGGTGGGGTTTCTTGAAGCCTAGTTCTCT
      220
-----insert pLM1-----
-----ORF pLM1-----
I D F L K K K N S E A Q A V I O G A L N A S E T T P K E L R I K R
CAAAACTCCTCAGATAGCATCTCAAGCCTCAACAGCATCACTAGCCATTCCAGCATCGGCAGCAGCAAGGATGCTGATGCCAAAAAGAAGAAAAAAGA
GTTTTGAGGAGTCTATCGTAGAGTTCGGAGTTGTCGTAGTGATCGGTAAGGTCGTAGCCGTCGTCGTTCTACGACTACGCTTTTCTCTTTTTTCTCT
      300
-----insert pLM1-----
-----ORF pLM1-----
C N S S O S I S S L N S I T S H S S I G S S K D A D A K K K K K L
GTTGGGTCTATGAGCTTCGAAGTTCTTCAACAAAGCGTTCAAGTATAAAAAAGGGGCCAAGTCAGCTTCCTCATACTCGGATATAGAGGAGATTGCTAC
CAACCCAGATACTCGAAGCTTCAAGGAAGTTGTTTCGAAGTCATATTTTTCCTCCGGGTCAGTCGAAGGAGTATGAGCCTATATCTCTCTAACGATG
      380
-----insert pLM1-----
-----ORF pLM1-----
S V V Y E L R S S F N K A F S I K K G P K S A S S Y S O I E E I A T
AGCCGACTCTTCAGCCCCCTCATCCCCAACTACAGCATGGTTCCACAGAGACTGCTTCACCCCTCATCAAGTCTCCACCTTGTCCTCCGTGGGCACT
TGGGCTGAGAAGTCGGGGGAGTAGGGGGTTTGATGTCGTACCAAGGTGTCCTGACGAAGTGGGAGGTAGTTCAGGAGGTGGAACAGGAGGCACCCGCTGA
      460
-----insert pLM1-----
-----ORF pLM1-----
P O S S A P S S P K L C H G S T E T A S P S I K S S T L S S V G T
GATGTCACCGAGGGCCCTGCTCAGCCAGCCCCACACTAGGCTGTTCCATGCAAAATGAGGAGGAGGAGCCAGAGAAGAAGGAGGTATCGGAGCTGCGCT
CTACASTGGCTCCCGGGACGAGTGGGTGCGGGGGTGATCCGACAAGGTACGTTTACTCCTCCTCCCGGTCCTTCTTCCATAGCCTCGACGCGA
      540
-----insert pLM1-----
-----ORF pLM1-----
D V T E G P A H P A P H T R L F H A N E E E E P E K K E V S E L R
CTGAGCTATGGGAGAAGGAAATGAAGCTTACAGACATCCCTTGGAGGGCCCTCAACTCTGCCCACTGGATCAGCTTCGGGAGACCATGCACAAACAT
GACTCGATACCTCTTCCTTTACTTCGAATGTCCTGAGGCGAACCCTCGGGAGTTGAGACGGGTGGTTGACCTAGTCGAAGCCCTCTGGTACGTGTTGTA
      620
-----insert pLM1-----
-----ORF pLM1-----
S E L V E K E N K L T C R L E A L N S A H Q L O Q L P E T I H N I

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GCAGTTGGAGGTGGACCTGCTGAAAGCAGAGAATGACCGACTGAAGGTAGCCCCAGGCCCTCATCAGGCTCCACTCCAGGSCAGGTCCCTGGATCATCT  
CGTCAACCTCCACCTGGACGACTTTCGTCTCTTACTGCGCTGACTTCCATCGGGGTCCGGGGAGTAGTCCGAGGTGAGGTCCCGTCCAGGACCTAGTAGA  
420  
-----insert pLM1-----  
-----ORF pLM1-----  
Q L E V D L L K A E N D R L K V A P G P S S G S T P G Q V D G S S  
GCATTATCTTCCCCACGCCGCTCCCTAGGCCCTGGCACTCACCCATTCCCTTCGCCCCAGTCTTGCAGACACAGACCTGTCACCCATGGA7GGCATCAGTA  
CGTAATAGAAGGGGTGCGGGGAGGGATCCGGACCGTGAGTGGGAAGGAAGCCGGGGTCAGAACGCTCTGTGCTGGACAGTGGGTACCTACCGTAGTCAT  
480  
-----insert pLM1-----  
-----ORF pLM1-----  
A L S S P R R S L G L A L T H S F G P S L A D T D L S P M D G I S  
CTTGTGGTCCAAAGGAGGAAGTGACCCCTCCGGGTGGTGGTGAGGATGCCCCCGCAGCACATCATCAAAGGGGACTTGAAGCAGCAGGAATTCTTCTGGG  
GAACACCAGGTTTCCTCCTTCACTGGGAGGCCACCACCACTCCTACGGGGGCGTCTGTAGTAGTTTCCCTGAACCTCGTCGCTCTAAGAAGGACCC  
540  
-----insert pLM1-----  
-----ORF pLM1-----  
T C G P K E E V T L R V V V R M P P Q H I I K G D L K Q Q E F F L G  
CTGTAGCAAGGTCAAGTGGAAAAGTTGACTGGAAGATGCTGGATGAAGCTGTTTTCAAGTGTTCAGGACTATATTTCTAAATGGACCCAGCCCTCTACC  
580  
GACATCGTTCCAGTCACCTTTTCAACTGACCTTCTACGACCTACTTCGACAAAAGGTTTCAAGTTTCTGATATAAAGATTTTACCTGGGTCGGAGATGG  
-----insert pLM1-----  
-----ORF pLM1-----  
C S K V S G K V D W K M L D E A V F Q V F K D Y I S K M D P A S T  
CTGGGACTAAGCACTGAGTCCATCCATGGCTACAGCATCAGCCACGTGAACAGAGTGTGGATGCAGAGCCCCCGAGATGCCTCCTTCCGTCGAGGTG  
620  
GACCTGATTCGTGACTCAGGTAGGTACCGA7GTCGTAGTCGGTGCACTTTCTCACAACCTACGTCCTGGGGGGCTCTACGGAGGAACGGCAGCTCCAC  
-----insert pLM1-----  
-----ORF pLM1-----  
L G L S T E S I H G Y S I S H V K R V L D A E P P E M P P C R R G  
TCAATAACATATCAGTCTCCCTCAAAGGTCTTAAGGAGAAATGCGTCGACACCCCTGGTGTTCAGACGCTGATCCCCAAGCCGATGATGCAGCACTACAT  
660  
AGTTATTTGTATAGTCAGAGGAGTTTCCAGACTTCTCTTTACGCAGCTGTGCGACCACAAGCTCTGCGACTAGGGGTTCCGGCTACTACGTCGTGATGTA  
-----insert pLM1-----  
-----ORF pLM1-----  
V Y N I S V S L K G L K E K C V D S L V F E T L I P K P M R C H Y I
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445CC TCCTGCTGAAGCACCGGCGCTCGTCTCTCGGCCCCAGCGGCACGGGCAAGACC TACCTGACCAATCGCTTGGCCGAGTACCTGGTGGAGCGC
TTCGGAGGACGACTTCGTGGCCGCGGAGCAGGAGAGCCCGGGGTGCGCGTGCCCGTTCTGGA TGGAC TGGTTAGCGAACC GGC TCATGGACCACCTCGCG
540
-----insert pLM1-----
-----ORF pLM1-----
S L L L K H R R L V L S G P S G T G K T Y L T N R L A E Y L V E R
TCTGGCCGTGAGGTCACAGAGGGCATCGTCAGCACCTTCAACATGCACCAGCAGTCTTGCAAGGATCTGCAACTGTATCTTCCAACTAGCCAACCAGA
AGACCGGCACCTCAGTGCTCCCGTAGCAGTCGTGGAAGTTGTACGTGGTCTGCAGAACGTTCC TAGACGTTGACATAGAAAGGTTGGATCGGTTGGTC
550
-----insert pLM1-----
-----ORF pLM1-----
S G R E V T E G I V S T F N M H Q Q S C K D L Q L Y L S N L A N G
TASACCGGGAACAGGAATTGGGGATGTGCCCTGGTGATTCTATTGGATGACCTGAGTGAAGCAGGCTCCATCAGTGAGTTGGTCAATGGGGCCCTCAC
ATCTGGCCCTTTGTCTTAACCCCTACACGGGGACCACTAAGATAACC TACTGGACTCAC TTCGTCCGAGGTAGTCAC TCAACCAGTTACCCCGGGAGTG
560
-----insert pLM1-----
-----ORF pLM1-----
I D R E T G I G D V P L V I L L D D L S E A G S I S E L V N G A L T
CTCAAGTATCATAAATGTCCTATATTATAGGTACCACCAATCAGCCTGTAAAAATGACACCCAACCATGGCTTGCACTTGAGCTTCAGGATGTTGACC
GACGTTTATAGTATTTACAGGGATATAATATCCATGGTGGTTAGTCGGACATTTT TACTGTGGGTTGGTACCGAACGTGAAC TCGAAGTCC TACAACCTGG
570
-----insert pLM1-----
-----ORF pLM1-----
C K Y H K C P Y I I G T T N Q P V K M T P N H G L H L S F R M L T
TCTCCAAACAGCTGGAGCCAGCCAATGGCTTCCTGGTTTCGTTACCTGAGGAGGAAGCTGGTAGAGTCAGACAGCGACATCAATGCCAACAGGAAGAGC
AAGAGGTTGTTGCACCTCGGTCGGTTACGGAAGGACCAAGCAATGGACTCC TCCTTCGACCATCTCAGTCTGTGCTGTAGTTACGGTTGTTCTTCTCG
580
-----insert pLM1-----
-----ORF pLM1-----
F S N N V E P A N G F L V R Y L R R K L V E S D S D I N A N K E E
TGGTTCGGGTGCTCGACTGGGTACCCAAAGCTGTGGTATCATCTCCACACCTTCCTTGAGAAGCACAGCACCTCAGACTTCCTCATCGGCCCTTGCTTCT
ACGAAGCCACAGAGCTGACCCATGGGTTTCGACACCATAGTAGAGGTGTGGAAGGAAC TCTTCGTGTCGTGGAGTCTGAAGGAGTAGCCGGGAACCAAGAA
590
-----insert pLM1-----
-----ORF pLM1-----
L L R V L D V V P K L V H L H T F L E K H S T S D F L I G P C F F

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[illegible]

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CATTCAAATATGTATCCGCTCATGAGACAATAACCCGTGATAAATGCTTCAATAATATTGAAAAAGGAAGAGTCCTGAGGGCGAAAGAACAGCTGTGGAA
GTAAGTTTATACATAGGCGAGTACTCTGTTATTGGGACTATTACGAAGTTATTATAACCTTTTCCTTCTCAGGACTCCGCCCTTCTTGGTCGACACCTT
H S N M Y P L M R Q . P . . M L Q . Y . K R K S P E A E R T S C G 7600

TGTGTGTCAGTTAGGGTGTGGAAAGTCCCCAGGCTCCCCAGCAGGCAGAAGTATGCAAAGCATGCATCTCAATTAGTCAGCAACCAGGTGTGGAAAGTCC
ACACACAGTCAATCCACACCTTTCAGGGGTCCGAGGGGTCTGTCCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTCCACACCTTTCAGG
M C V S . G V E S P Q A P Q Q A E V C K A C I S I S Q Q P G V E S P 7700

CCAGGCTCCCCAGCAGGCAGAAGTATGCAAAGCATGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCCATCCCGCCCTAACTCCGC
GGTCCGAGGGGTCTGCTCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTATCAGGGCGGGGATTGAGGCGGGTAGGGCGGGGATTGAGGCS
Q A P Q Q A E V C K A C I S I S Q Q P . S R P . L R P S R P . L R 7800

CCAGTTCGCCCCATTCTCCGCCCCATGGCTGACTAATTTTTTTTATTTATGCGAGGCGGAGGCGCCTCGGCTCTGAGCTATTCCAGAAGTAGTGAGG
GGTCAAGGCGGGTAAGAGGCGGGTACCGACTGATTAAAAAAATAAATACGTCCTCCGGCTCCGCGGAGCGGAGACTCGATAAGGCTTCATCACTCC
P V P P I L R P H A D . F F L F M Q R P R P R P L S Y S R S S E 7900

AGGCTTTTTTGGAGGCTAGGCTTTTGGAAAGATCGATCAAGAGACAGGATGAGGATCGTTTCGATGATTGAACAAGATGGATTGCACGCAGGTTCTCC
TCCGAAAAAACCTCCGGATCCGAAACGTTTCTAGCTAGTTCTCTGTCTACTCTAGCAAAGCGTACTAAGTTGTTCTACCTAACGTGCGTCCAAGAGG
E A F L E A . A F A K I D Q E T G . G S F R M I E Q D G L H A G S P 9000

GGCGCTTGGGTGGAGAGGCTATTCCGCTATGACTGGGCACAACAGACAATCGGCTGCTCTGATGCCCGCTGTTCCGGCTGTGAGCGCAGGGGCGCCCG
CCGGCGAACCCACCCTCCGATAAGCCGATACTGACCCGTGTTGCTGTTAGCCGACGAGACTACGGCGGCACAAGGCCGACAGTCGCGTCCCCGGGGC
A A V V E R L F G Y D V A Q Q T I G C S D A A V F R L S A Q G R P 9100

GTTCTTTTGTCAAGACCGACCTGTCCGGTGCCCTGAATGAACTGCAAGACGAGGCGAGCGGGCTATCGTGGCTGGCCACGACGGGCGTTCTTGCACG
CAAGAAAAACAGTTCTGGCTGGACAGCCACGGGACTTACTTGACGTTCTGCTCCGTCGCGCCGATAGCACCGACCGGTGCTGCCCGCAAGGAACCGGCTC
V L F V K T D L S G A L N E L Q D E A A R L S V L A T T G V P C A 9200

CCTGCTCGACGTTGTCACTGAAGCAGGAGGACTGSGTGCTATTGGGCGAAGTGCCGGGGCAGGATCTCCTGTCTCTCACCTTGCTCCGCGAGAA
GACACGAGCTGCAACAGTGACTTCGCTTCCCTGACCGACGATAACCCGCTTACGCGCCCGTCTTAGAGGACAGTAGAGTGGAACGAGGACGGCTCTT
A V L D V V T E A G R D V L L L G E V P G Q D L L S S H L A P A E 9300

AGTATCCATCATGGCTGATGCAATGCGGCGGCTGATACGCTTGATCCGGCTACCTGCCCCATTCGACCACCAAGCGAAACATCGCATCGAGCGAGCAGC
TCATAGGTAGTACCGACTACGTTACGCGCGGAGCTATGCGAATAGGCCGATGGACGGGTAAGCTGGTGGTTGCTTTGTAGCGTAGCTCGCTCTGTGCA
V S I M A D A M R R L H T L D P A T C P F D H Q A K H R I E R A R 9400

ACTCGGATGGAAGCCGGCTTTGTGCTATGAGGATGATGAGGACGAGCATCAGGGGCTCGCGCCAGCCGAACGTTTCGCCAGGCTCAAGGCGAGCATGC
TGAGGCTACCTTCGGCCAGAACAGCTAGTCCCTACAGAGCTGCTTCTGCTAGTCCCCGAGCGCGGTCGGCTTGACAAGCGGTCCGAGTTCCGCTCTGTAC
R M E A G L V C Q D D L D E E H Q G L A P A E L F A R L K A S P 9500

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CCGACGGCGAGGATCTCGTGGTACCCATGGCGATGCCTGCTTGCCGAATATCATGGTGGAAAAATGGCCGCTTTCTGGATTTCATCGACTGTGGCCGGCT
GGCTGCCGCTCCTAGAGCAGCAC TGGGTACCGCTACGGACGAACGGCTTATAGTACCACCTTTTACCGGCGAAAAGACCTAAGTAGCTGACACCGGCCGA 9600

Kan/Neo

P D G E D L V V T H G D A C L P N I M V E N G R F S G F I D C G R L

GGGTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCGCTTCTCTGCTCTTACGGT
CCACACCGCCTGGCGATAGTCC TGTATCGCAACCGATGGGCACTATAACGACTTCTCGAACCGCCGCTTACCCGACTGGCGAAGGAGCACGAAATGCCA 9700

Kan/Neo

G V A D R Y Q D I A L A T R D I A E E L G G E V A D R F L V L Y G

ATCGCCGCTCCCGATTGCGACGCGCATCGCCTTCTATCGCCTTCTTGACGAGTTCTTCTGAGCGGGACTCTGGGGTTCGAAATGACCGACCAAGCGACGCG
TAGCGGCGAGGGCTAAGCGTCGCGTAGCGGAAGATAGCGGAAGAAGCTGCTCAAGAAGACTCGCCCTGAGACCCCAAGCTTTACTGGCTGGTTGCTGCGG 9800

Kan/Neo

I A A P D S Q R I A F Y R L L D E F F . A G L W G S K . P T K R R

CAACCTGCCATCAGGAGATTTCGATTCCACCGCCGCTTCTATGAAAGGTTGGGCTTCGGAATCGTTTTCCGGGACGCCGGCTGATGATCCTCCAGCGC
GTTGGAGGTTAGTGCTCTAAAGCTAAGGTGGCGGCGGAAGATACCTTCAACCCGAAGCCTTAGCAAAAGGCCCTGCGGCCGACCTACTAGGAGGTGCGG 9900

P T C H H E I S I P P P P S M K G V A S E S F S G T P A G . S S S A

GGGATCTCATGCTGGAGTCTTCGCCCCACCTAGGGGGAGGCTAACTGAAACACGGAAGGAGACAATACCGGAAGGAACCCGCGCTATGACGGCAATAA
CCCCTAGAGTACGACCTCAAGAAGCGGGTGGGATCCCCCTCCGATTGACTTTGTGCTTCTCTGTTATGGCTTCTCTGGGCGCGATAGTCCGCTTATT 9000

G I S C V S S S P T L G G G . L K H G R R Q Y R K E P A L . R Q .

AAAGACASAATAAAACGCAGCGGTGTTGGGTGCTTGTTCATAAACCGGGGTTGCGTCCAGGGCTGGCACCTCTGTCGATACCCACCGAGACCCCATTS
TTTCTGCTCTATTTGCGTGCCACAACCCAGCAACAAGTATTTGCGCCCCAAGCCAGGCTCCCGACCGTGAGACAGCTATGGGGTGGCTCTGGGGTAAC 9100

X D R I K R T V L G R L F I N A G F G P R A G T L S I P H R D P I

GGGCCAATACGCCCCGCTTCTTCTTTTCCCCACCCACCCCAAGTTGCGGTGAAGGCCAGGGCTCGCAGCCAACGTCGGGGCGGCGAGCCCTGCG
CCCGGTTATGCGGGCGCAAGAAGGAAAAGGGTGGGGTGGGGGTTCAAGCCCACTTCCGGGTCCCGAGCGTGGGTTGCAAGCCCGCGCTCGGGACGG 9200

G A N T P A F L P F P H P T P Q V R V K A O G S Q P T S G R Q A L F

ATAGCCTCAGGTTACTCATATATACTTTAGATTGATTTAAACTTCATTTTAAATTTAAAGGATCTAGGTGAAGATCCTTTTGTATAATCTCATGACCA
TATCGGAGTCCAATGAGTATATGAAATCTAACTAAATTTGAAGTAAAAATTAATTTTCTAGATCCACTTCTAGGAAAAATATTAGAGTACTGGT 9300

P Q V T H I Y F R L I . N F I F N L K G S R . R S F L I I S . F

AAATCCCTTAACGTGAGTTTTCGTTCCACTGAGCGTCAGACCCCGTAGAAAAGATCAAGGATCTTCTTGAGATCCTTTTCTTCGCGTAATCTGCTG
TTTAGGGAATTGCACITCAAAGCAAGGTGACTCGCACTCTGGGGCATCTTTCTAGTTTCTAGAGAAGCTCTAGGAAAAAAGAGCGCGCATTAGACGAC 9400

pUC ori

K S L N V S F R S T E R Q T P . K R S K D L L E I L F F C A . S A

CTTGCAAAACAAAAAACCCGCTACCAAGCGGTGGTTTGTGCGGATCAAGAGCTACCAACTCTTTTCCGAAGGTAACGGCTTCAGCAGAGCGGAG
GAACGTTGTTTTTTGGTGGCGATGGTGGCCACCAACCAACCGGCTAGTTCTCGATGGTTGAGAAAAAGGCTTCATTGACCGAAGTCTCTCGCGTC 9500

pUC ori

A C K Q K N H R Y Q P V F V C R I K S Y O L F F R R . L A S A E R F

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ATACCAAATACTGTCTTC TAGTGTAGCCGTAGTTAGGCCACCACTTCAAGAACTCTGTAGCACCGCCTACATACCTCGCTCTGCTAATCCTGTTACCA3
TATGGTTTATGACAGGAAGATCACATCGGCATCAATCCGGTGGTGAAGTTC TTGAGACATCGTGGCGGATGTATGGAGCGAGACGATTAGGACAATGGTC 960

pUC ori
Y Q I L S F . C S R S . A T T S R T L . H R L H T S L C . S C Y Q

TGGCTGCTGCCAGTGGCGATAAGTCGTGCTTACCGGGTGGACTCAAGACGATAGTTACCGGATAAGGCGCAGCGGTGGGGCTGAACGGGGGGTTCGTS
ACCGACGACGGTCACCGCTATTCAGCACAGAATGGCCCAACCTGAGTTCTGCTATCAATGGCCTATTCCGCGTCGCCAGCCGAC TTGCCCCCAAGCAC 970

pUC ori
V L L P V A I S R V L P G V T Q D D S Y R I R R S G R A E R G V R

CACACAGCCCAGCTTGGAGCGAAGCAGCTACACCGAACTGAGATACCTACAGCGTGAGCTATGAGAAAGCGCCACGCTTCCGAAGGGAGAAAGCGGAC
GTGTGTCGGGTCGAACCTCGCTTGCTGGATGTGGCTTGACTCTATGGATGTGCGACTCGATACTCTTTCGCGGTGCGAAGGGCTTCCTCTTTCCGCC TG 980

pUC ori
A H S P A V S E R P T P N . D T Y S V S Y E K A P R F P K G E R R T

AGGTATCCGGTAAGCGGCAGGGTCGGAACAGGAGAGCGCACGAGGGAGCTTCCAGGGGGAACGCCTGGTATCTTTATAGTCC TGTCGGGTTTCGCCACC
TCCATAGGCCATTTCGCGTCCAGCCTTGCTCTCGCGTGC TCCCTCGAAGGTCCCCCTTTGCGGACCATAGAAATATCAGGACAGCCCAAAGCGGTGG 990

pUC ori
G I R . A A G S E Q E S A R G S F Q G E T P G I F I V L S G F A T

TCTGACTTGAGCGTCGATTTTGTGATGCTCGTCAGGGGGGCGGAGCCTATGGAAAAACGCCAGCAACGCGGCCTTTTACGGTTCTTGCCCTTTTGCTG
AGACTGAACTCGCAGCTAAAAACACTACGAGCAGTCCCCCGCCTCGGATACCTTTTTCGGTTCGTGCGCCGGAATAATGCCAAGGACCGGAAAAACGAC 1000

pUC ori
S D L S V D F C D A R Q G G G A Y G K T P A T R P F Y G S V P F A

GCCTTTTGCTCACATGTTCTTTCTGCGTTATCCCTGATTCTGTGGATAACCGTAT TACCGCCATGCAT 10070
CGGAAAACGAGTGTACAAGAAAGGACGCAATAGGGGACTAAGACACCTATTGGCATAATGGCGGTACGTA
G L L L T C S F L R Y P L I L V I T V L P P C I

SEQUENCE ID No. 8.

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fig 30 pEGFP72 (1 > 9697) Site and Sequence

Enzymes: 72 of 148 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

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16p

Bgl I

TAGTTATTAAATAGTAATCAATTACGGGGTCATTAGTTTCATAGCCCATATATGGAGTTCGCGTTACATAAATTACGGTAAATGGCCCGCTGGCTGACCG 100

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L L I V I N Y G V I S S . P I Y G V P R Y I T Y G K V P A V L T

Aat II

CCCAACGACCCCGCCCATTTGACGTCAATAATGACGTATGTTCCCATAGTAACGCCAATAGGGACTTTCCATTGACGTCAATGGGTGGAGTATTTACGGT 200

GGGTTGC TGGGGCGGTAAC TGCAGTTATTACTGCATACAAGGGTATCATTGCGGTATCCCTGAAAGGTAAGTGCAGTTACCCACCTCATAAATGCCA

A Q R P P P I D V N N D V C S H S N A N R D F P L T S M G G V F T V

Bgl I

AAACTGCCCACTTGGCAGTACATCAAGTGTATCATATGCCAAGTACGCCCCCTATTGACGTCAATGACGGTAAATGGCCCGCTGGCATTATGCCCAGTA 300

TTTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTTCATGCGGGGATAACTGCAGTTACTGCCATTACCGGGCGGACCGTAATACGGGTCAT

N C P L G S T S S V S Y A K Y A P Y . R Q . R . M A R L A L C P V

SnaB I

CATGACCTTATGGGACTTTTCTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGATGCGGTTTTTGGCAGTACATCAATGGGCGTGGG 400

GTA CTGGAATACCTGAAAGGATGAACCGTCATGTAGATGCATAATCAGTAGCGATAATGGTACCCTACGCCAAAACCGTCATGTAGTTACCCGACCT

H D L H G L S Y L A V H L R I S H R Y Y H G D A V L A V H Q V A V

Aat II

TACGGGTTTGACTCAGCGGATTTTCAAGTCTCCACCCATTGACGTCAATGGGAGTTTGTTTTGGCACCAAAATCAACGGGACTTTCCAAAATGTCGTG 500

ATGCCAAACTGAGTGGCCCTAAAGGTTGAGAGTGGGGTAAC TGCAGTTACCCTCAAAACAAACCGTGGTTTTAGTTGCCCTGAAAGGTTTTACAGCAT

I A V . L T S I S K S P P H . R Q V E F V L A P K S T G L S K M S .

Nhe I

ACAAC TCCGCCCCATTGACGCAAAATGGCGGTAGGCGTGTACGGTGGGAGGTCTATATAAGCAGAGCTGGTTTAGTGAACCGTCAGATCCCTAGCGCTG 600

TGTTGAGGCGGGGTAAC TGGGTTTACCGCCATCGGCACATGCCACCTCCAGATATATTCGTCTCGACCAAAATCACTTGGCAGTCTAGGCGATCGCGAT

Q L R P I D A N G R . A C T V G G L Y K Q S V F S E P S D P L A L

Nco I

CGGTCGCCACCATGTTGAGCAAGGCGGAGGAGCTGTTCACGGGGTGGTGCCCATCTGGTTCGAGCTGGACGGCGACGTAAACGGGCCACAAATTCAGCG 700

GGGCAGCGGTGGTACCACTGGTTCCCGCTCCCTGCAAAAGTGGCCCCACACGGGTAGGACCAGCTCGACCTGCCGCTGCATTTGCCGGTGTCAAGTCGG

eGFP.C.e.unc53

F V A T M V S K G E E L F T G V V P I L V E L D G D V H G H K F S

Nco I

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V S G E G E S D A T Y S < L T L K F I C T T G K L P V P V P F L V T

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KspI

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1300

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BspM II Bgl II

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1400

eGFP.C.e.unc53

C.e.unc53

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Nru I EcoR I

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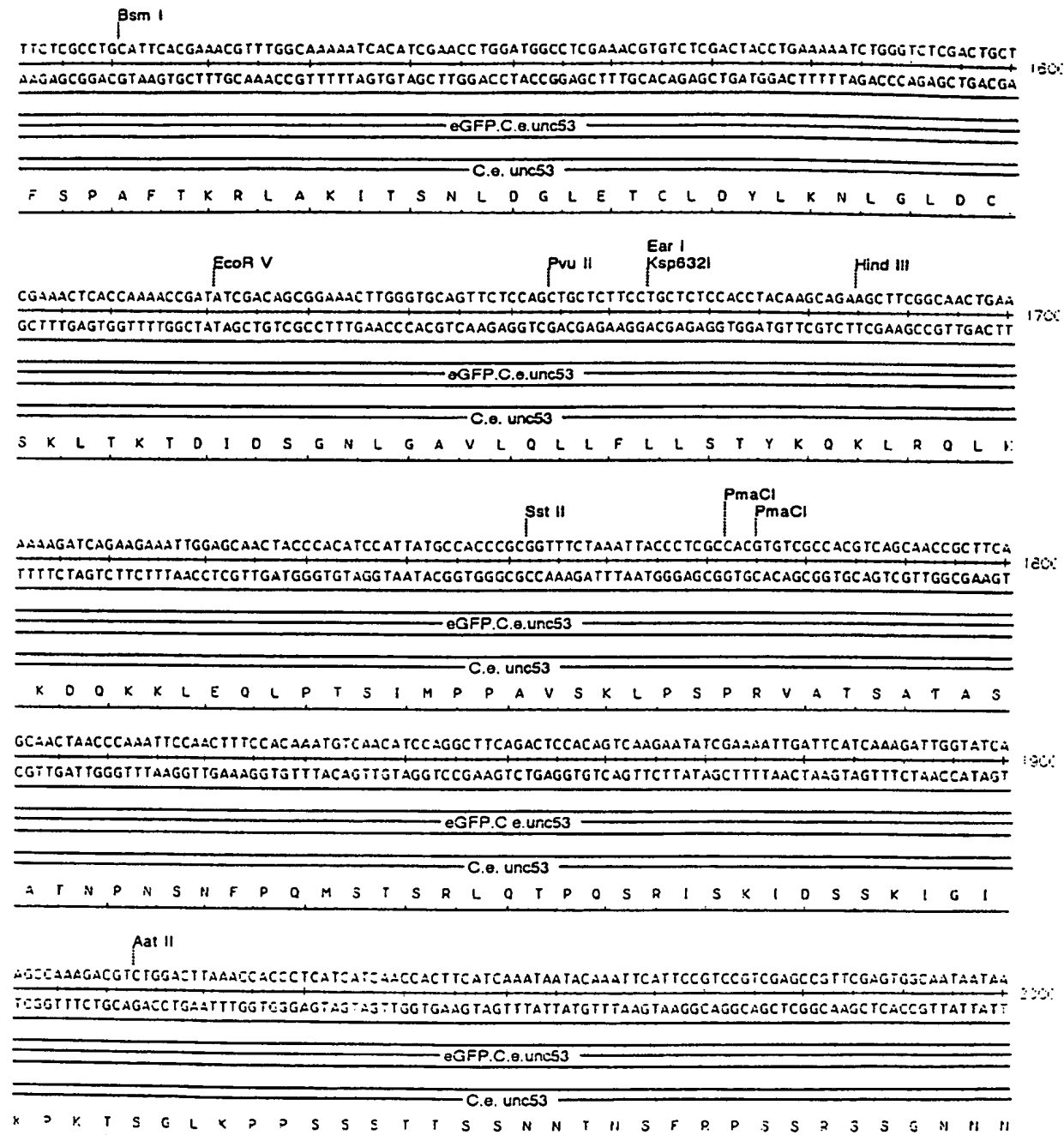
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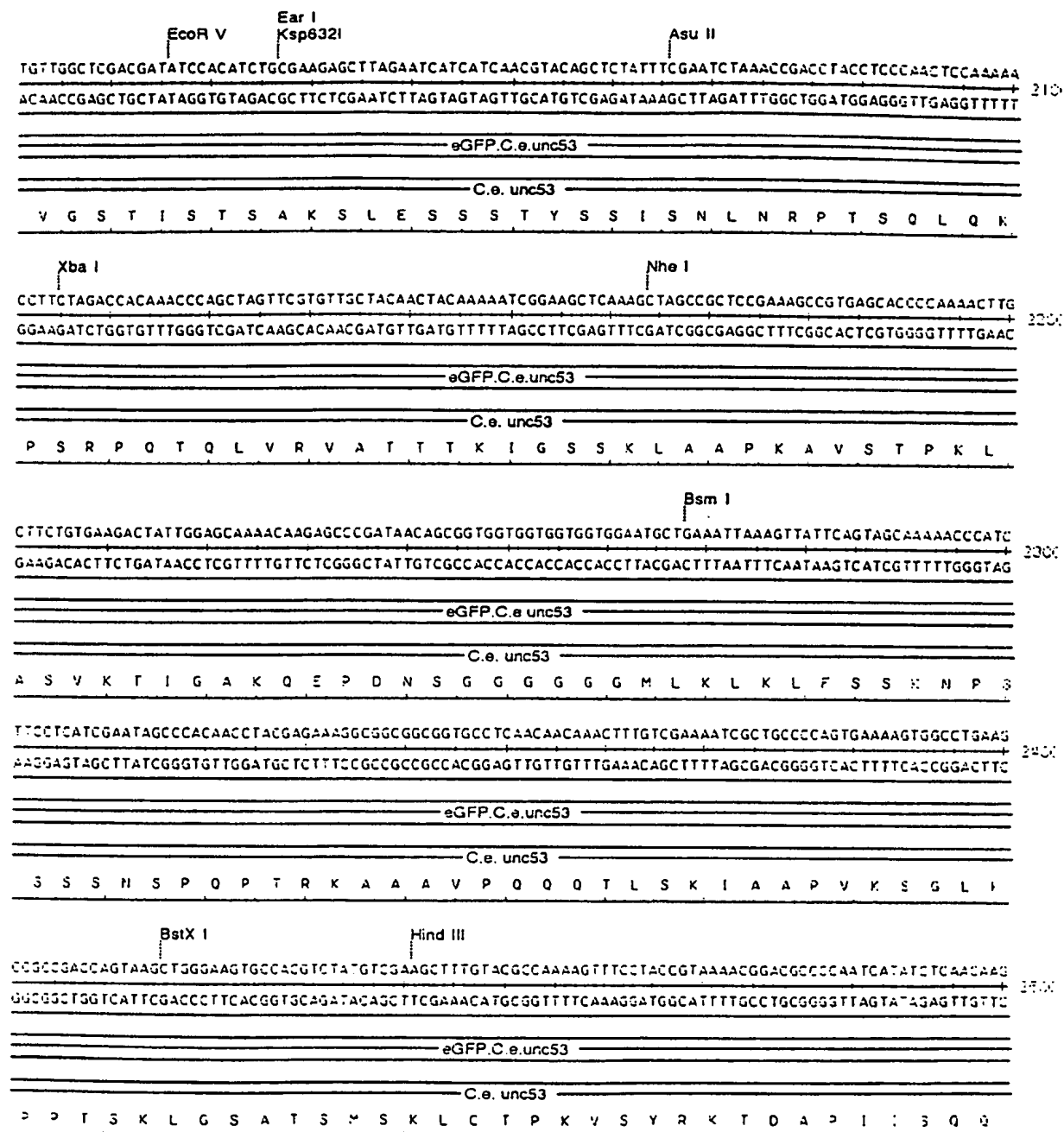
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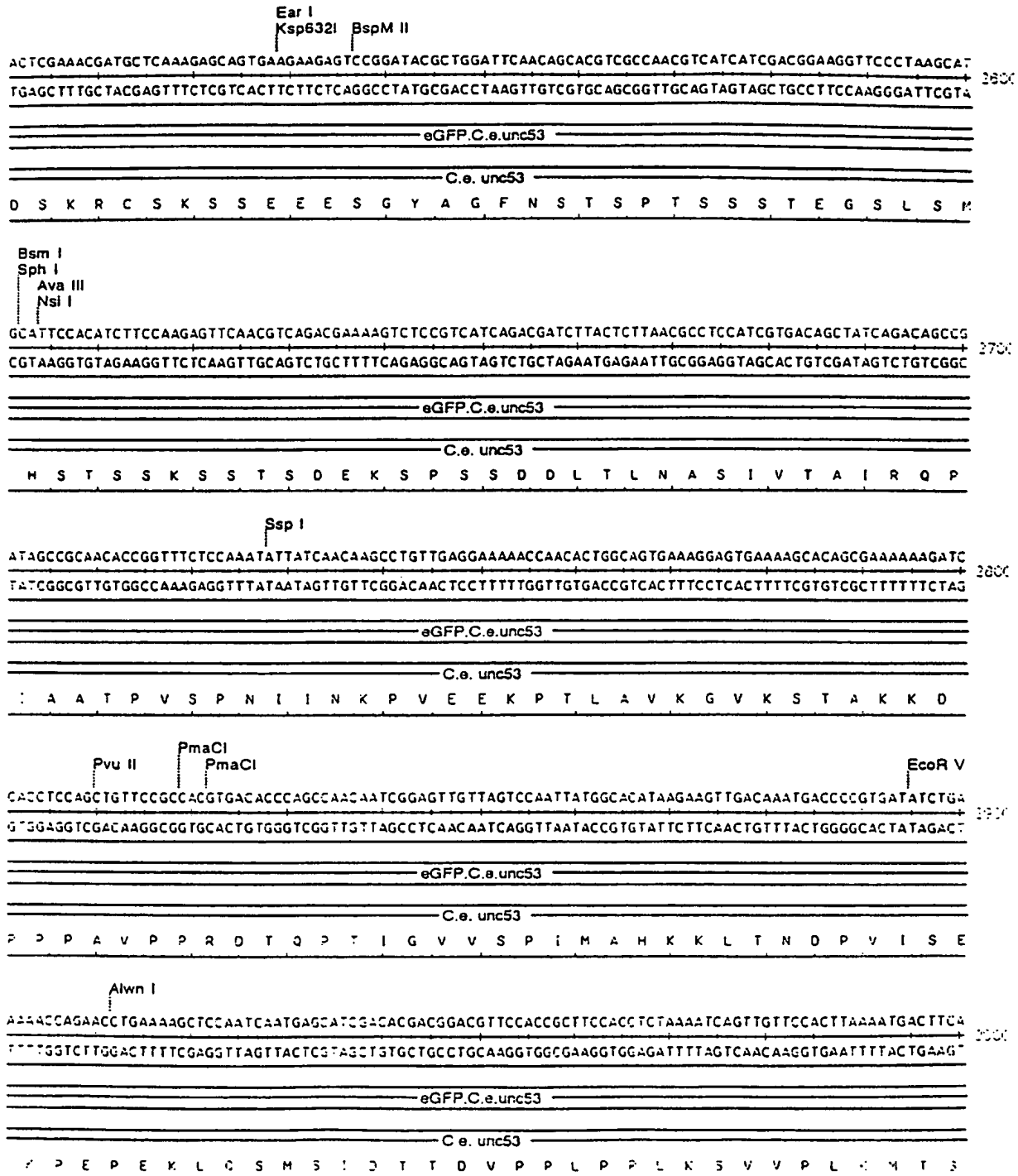
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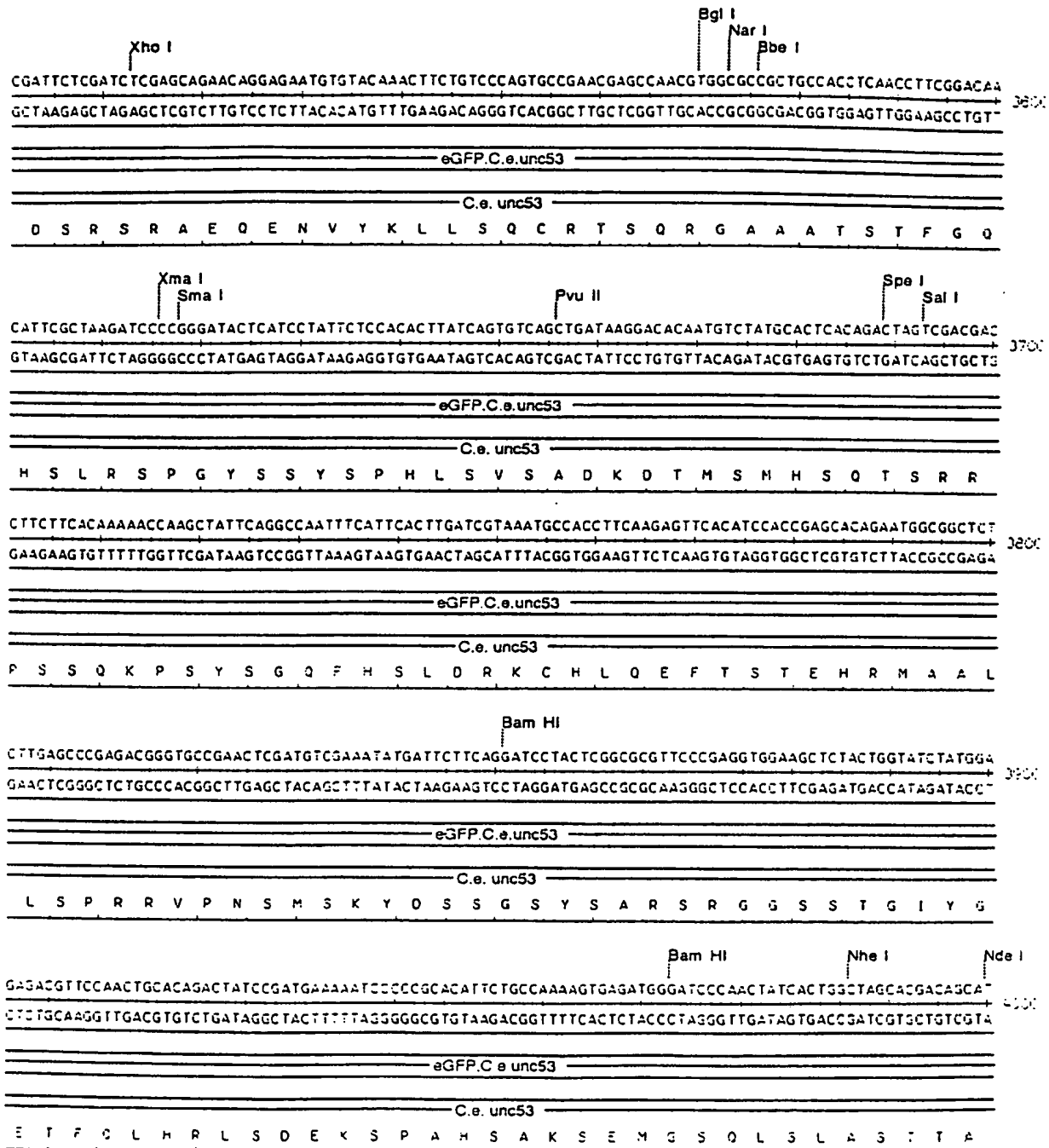
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TAGGCTGTTGGTGGTTGCAATGCTACAAGAAGATTTGTTCTTTTAGTGTAGCGGACAGTTCAGCAAACTATAC TCGTCAGCAGGCGCAGACTTCTGA
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-----C.e. unc53-----
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-----C.e. unc53-----
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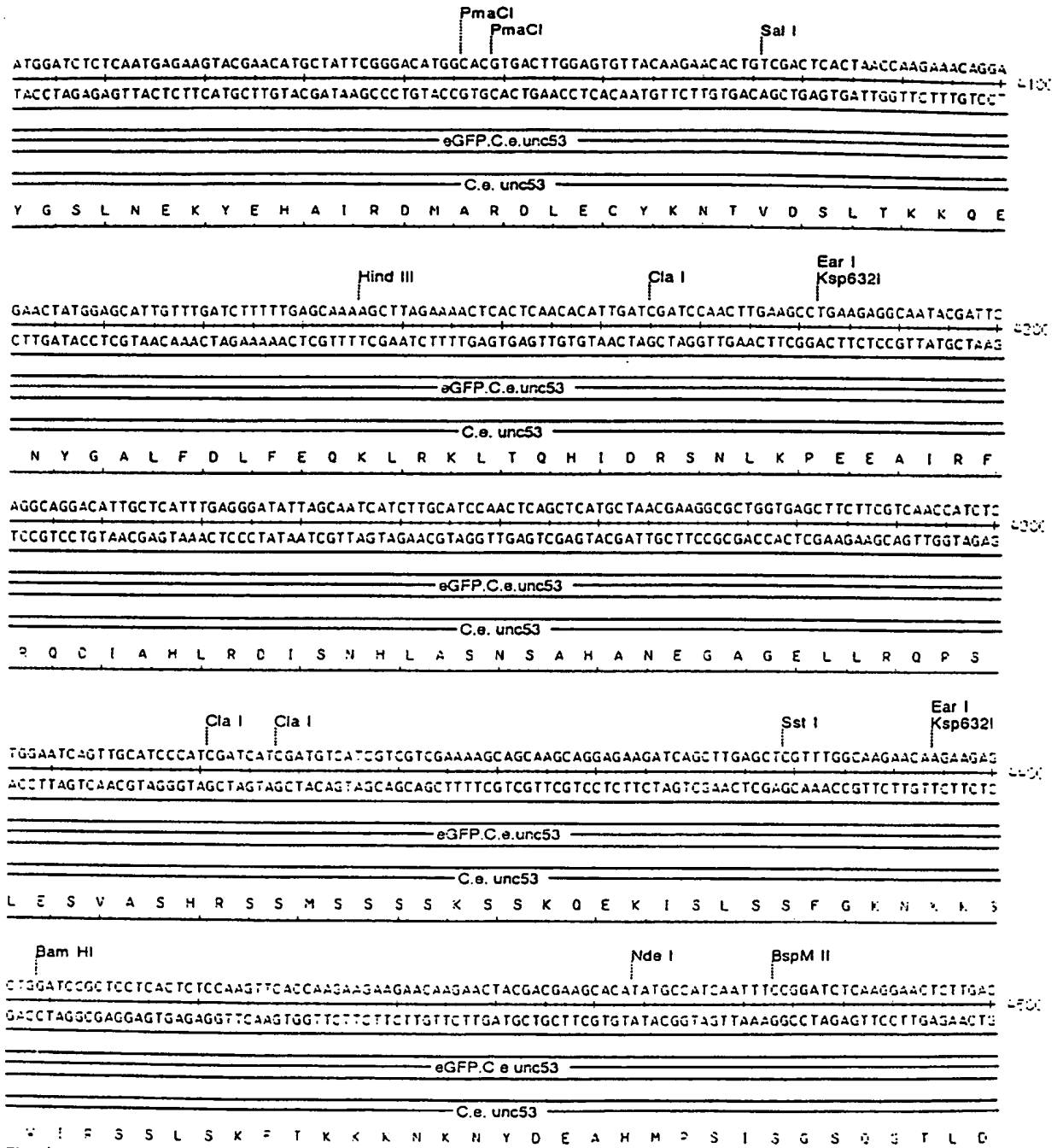
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----- C.e.unc53 -----
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KspI BsrI Asu II
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Pvu I Hpa I EcoR V
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Cla I
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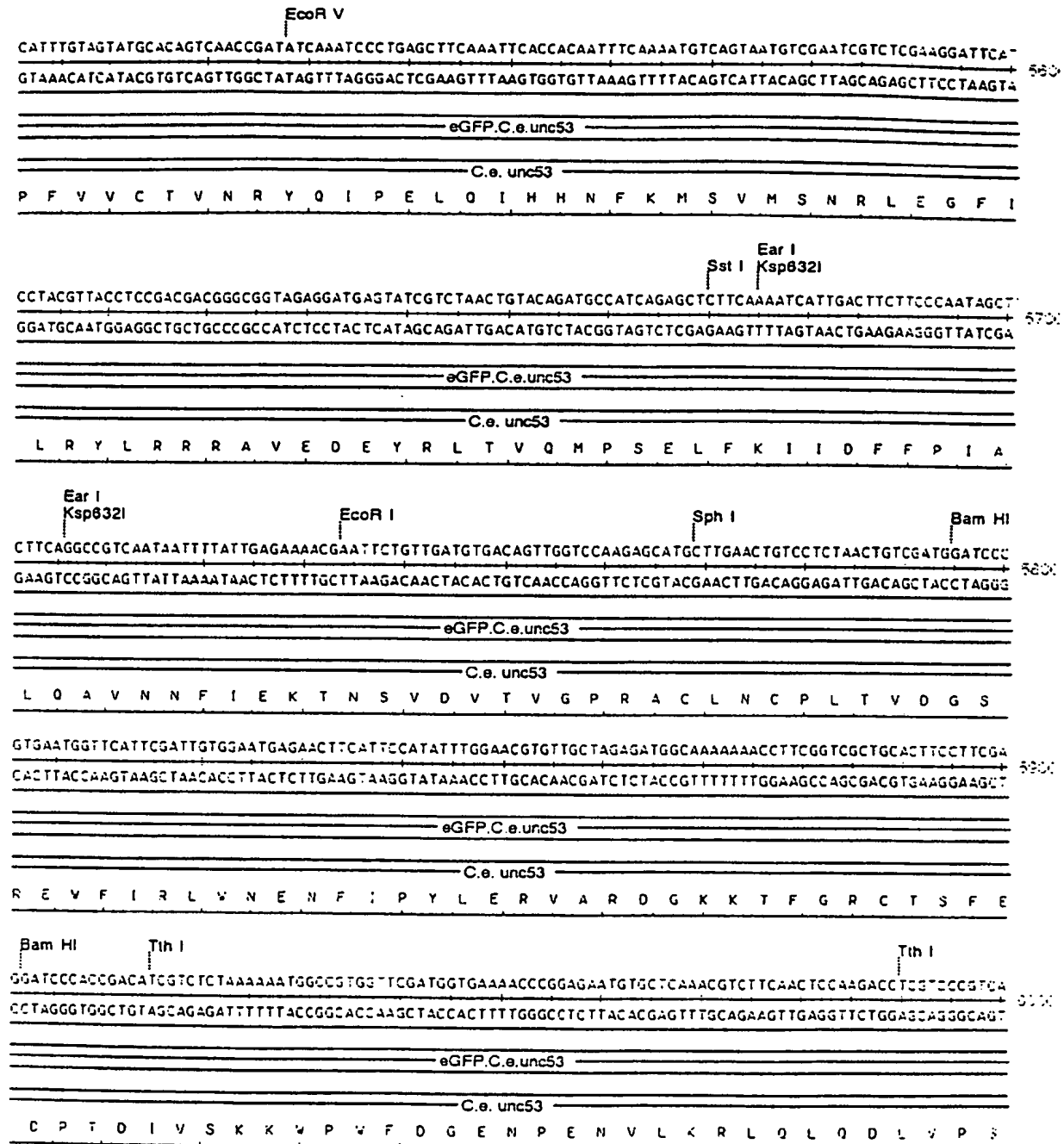
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Mlu I
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C.e. unc53
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C.e. unc53
T T I R M F M H G A A O S R V D S L V L D M L L P K Q M I L Q L V
Aat II Bsr I Bsr I Asu II
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Bsm I Xmn I Bgl II
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C.e. unc53
N O S E D S I V N I S I P E N N K E E L L Q V E R R L E K I L R S
Ava III Nsi I Xba I
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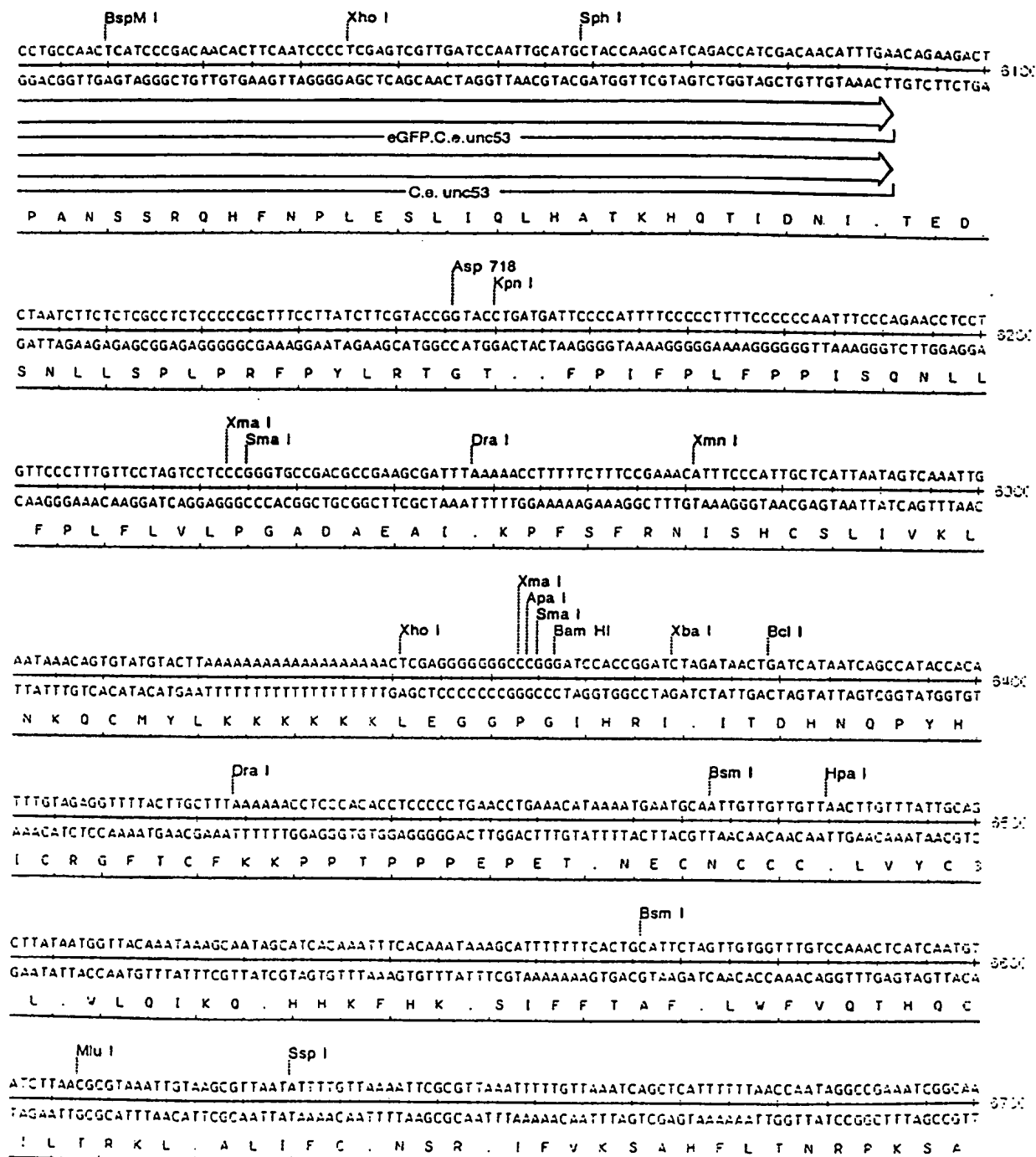
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Bsr I

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K S L I N Q K N R P R . G . V L F Q F G T R V H Y . R T V T P T S I 6800

Dra III

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Nae I

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L K G A P D L E L D G E S R R T V R E R K G R K R K E R A L G R V 7000

Ksp I

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Q V . R S R C A . P P H P P R L M R R Y R A R Q V A L F G E M C A E 7100

BspH I

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P L F V Y F S K Y I Q I C I R S . D N N P D K C F N N I E K G R V 7200

Sph I
Ava III
Nsi I

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Sph I
Ava III
Nsi I

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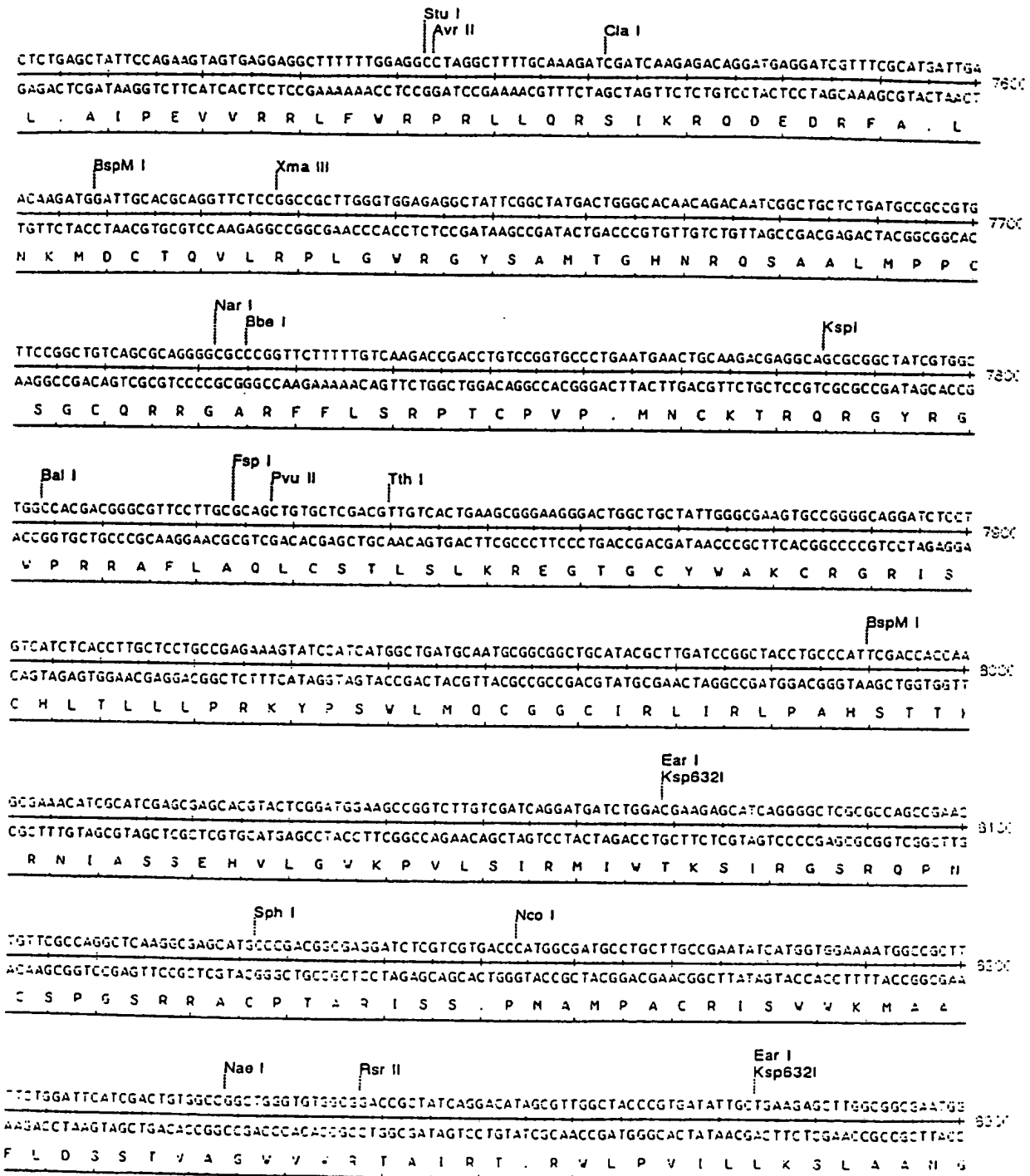
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Bgl I
Sfi I

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Asu II BspM I
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Nae I KspI Avr II
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KspI
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AlwI OxaN I Dra I Dra I
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BspH I
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BsrI
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A V G L N G G F V H T A Q L G A N D L H R T E I P T A . A M R K R 9400

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H A S R R E K G G Q V S G K R Q G R N R R A H E G A S R G K R L V S 9500

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L . S C R V S P P L T . A S I F V M L V R G A E P M E K R Q Q R G 9600

Ava III
Nsi I

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SEQUENCE I.D. No 9

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SEQUENCE ID NO 10

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fig 13 pCB201 (1 > 5082) Site and Sequence

Enzymes : 100 of 148 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

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P Y . R Q . R . N A R L A L C P V H D L M G L S Y L A V H L R I S H

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T7 promoter priming site

ProBond binding domain

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fig 13 pCB201 (1 > 5082) Site and Sequence

Page

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pCB201 insert = U4
U4 ORF
D D D K V P R I H M P P C R R G V N N I S V S L K G L K E K C Y D
GCCTGGTGTTCGAGACGCTGATCCCAAGCCGATGATGCAGCACTACATAAGCCTCCTGCTGAAGCACCAGCGCCTCGTCTCTCGGGCCCCAGCGGCAC
CGGACCACAAGCTCTGCGACTAGGGGTTCCGGCTACTACGTCGTGATGTATTCGGAGGACGACTTCGTGGCCCGCGGAGCAGGAGAGCCCCGGGGTCGCCGTG
pCB201 insert = U4
U4 ORF
S L V F E T L I P K P M M O H Y I S L L L K H R R L V L S G P S G T
GGGCAAGACCTACCTGACCAATCGCTTGCCCGAGTACCTGGTGGAGCGCTCTGGCCGTGAGGTCACAGAGGGCATCGTCAGCACCTTCAACATGCACCA
CCCGTTCTGGATGGACTGGTTAGCGAACCGGCTCATGGACCACCTCGCGAGACCGGCACTCCAGTGTCTCCCGTAGCAGTCGTGGAAGTTGTACGTGGT
pCB201 insert = U4
U4 ORF
G K T Y L T N R L A E Y L V E R S G R E V T E G I V S T F N M H Q
CACTCTTGAAGGATCTGCAACTGTATCTTTCCAACCTAGCCAACCAGATAGACCGGGAACAGGAATTGGGGATGTGCCCTTGGTGATTCATTGSA
GTCAAGACGTTCTAGACGTTGACATAGAAAGGTTGGATCGGTTGGTCTATCTGGCCCTTTGTCTTAACCCCTACACGGGGACCACAAAGATAACCTAC
pCB201 insert = U4
U4 ORF
V S C K O L O L Y L S N L A N O I D R E T G I G D V P L V I L L D
ACCTGAGTGAAGCAGGCTCCATCAGTGAGTTGGTCAATGGGGCCCTCACCTGCAAGTATCATAAATGTCCCTATATTATAGGTACCACCAATCAGCCTG
TGGACTCAGCTTCTGTCGAGGTAGTCACTCAACCACTTACCCCGGGAGTGGAGGTTCAATGATTTACAGGGATATAATATCCATGGTGGTATGTCGAC
pCB201 insert = U4
U4 ORF
D L S E A G S I S E L V A G A L T C K Y H K C P Y I I G T T N Q P
AAAAATGACACCCAACCATGGCTTGCACCTTGAGCTTCAAGGATGTGACCTTCTCCAACCAAGTGGAGCCAGCCAATGGCTTCTGGTTGGTTACCTGAG
TTTCTACTGTGGGTGGTACCGAACGTGAACCTCAAGTCTTACAACCTGGAAGAGGTTGTGACCTCGGTTCGGTTACCGAAGGACCAAGCAATGGACTC
pCB201 insert = U4
U4 ORF
K N T P N H G L H L S F R M L T F S N H V E P A N G F L V R Y L F

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AGGAAGCTGGTAGAGTCAGACAGCGACATCAATGCCAACAAGGAAGAGCTGCTTCGGGTGCTCGACTGGGTACCCAAGCTGTGGTATCATCTCCACACCT
TCCCTCGACCATCTCAGTCTGTCGCTGTAGTTACGGTTGTTCTCTCGACGAAGCCACGAGCTGACCCATGGGTTGACACCATAGTAGAGGTGTGGG

pCB201 insert = U4

U4 ORF

R K L V E S D S D I N A N K E E L L R V L D V V P K L V Y H L H T

TCCTTGAGAAGCACAGCACCTCAGACTTCCTCATCGGCCCTTGCTTTCTTGTGCTGTCCATTGGCATTGAGGACTTCCGGACCTGGTTCATTGACCT
AGGAACCTCTTCGTGCTGTGGAGTCTGAAGGAGTAGCCGGGAACGAAGAAAGACAGCACAGGGTAACCGTAACCTCTGAAGGCCGACCAAGTAACGGG

pCB201 insert = U4

U4 ORF

F L E K H S T S D F L I G P C F F L S C P I G I E D F R T W F I D L

GTGGAACACTCTATCATTCCCTATCTACAGGAAGGAGCCAAGGATGGGATAAAGGTCCATGGACAGAAAGCTGCTTGGGAGGACCCAGTGGAAATGGGT
CACCTTGTGAGATAGTAAGGGATAGATGTCTTCTCGGTTCTTACCCTATTTCAGGTACCTGTCTTTCGACGAACCTCTCGGGTACCTTACCCAG

pCB201 insert = U4

U4 ORF

V N N S I I P Y L Q E G A K D G I K V H G Q K A A W E D P V E W V

CGGGACACACTTCCCTGGCCATCAGCCCAACAAGACCAATCAAAGCTGTACCACCTGCCCCACCCACCGTGGGCCCTCACAGCATTGCCCTACCTCCCG
GCCCTGTGTGAAGGGACCGGTAGTCGGGTTGTCTGGTTAGTTTCGACATGTTGGACGGGGGTGGGTGGCACCCGGGAGTGTCTGAACGGAGTGGAGGGG

pCB201 insert = U4

U4 ORF

R D T L P V P S A Q Q D Q S K L Y H L P P P T V G P H S I A S P P

AGGATAGGACAGTCAAGACAGCACCCCAAGTTCTCTGGACTCAGATCCTCTGATGGCCATGCTGCTGAAACTTCAAGAAGCTGCCAATACATTGAGTC
TCCTATCTTGTGAGTTTCTGCTGTGGGTTCAAGAGACC TGAGTCTAGGAGACTACCGGTACGACGACTTTGAAGTTCTTCGACGTTGATGTAACTGAC

pCB201 insert = U4

U4 ORF

E D R T V K D S T P S S L D S D P L M A M L L K L Q E A A H Y I E S

TCCAGATGGAGAAACCATCCTGGACCCCAACCTTCAGGCAACACTTAAAGGTTTCGGCAATCACTGTACCCCCGGACAGCAGAACGCTGGCATCAGCTA
AGGTCATGCTCTTGTGAGGACCTGGGTTGGAAGTCCGTTGTGAAATTCCEAAGCCGTTAGTGACAGTGGGGGCCGTGCTGCTTTGCGACCGTATGCGA

pCB201 insert = U4

U4 ORF

P D F E T I L C P N L Q A T L G F G N H C H P R T A E R V H Q L

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TCTTAGCTCCTCCTCTCCCTCTCTCTTTTCAGAGCACTGGCTCTCCAGCCCCAGGAGGAGAACAGGAGGGAGGAGAGATGAAAGAGGAGGGACAGGT
AGAATCGAGGAGGAGAGGGGAGAGGAGAAAGTCTCGTGACCGAGAGGTGGGGTCTCTCTTTGTCTCCCTCCTCCTCTACTTTCTCTCCCTGTCCAA 2300

pCB201 insert = U4

S L L L S P L L F Q S T G S P A P G G E Q E G G G D E R G G T G

CTTGGTGCTGTACCTTTGAGAAC TTCTAGGAAGGAATGGTGGGGTGGCGTTTGGGAAC TTGTGCCCCCTAAACACATTTACTGGCCTCCTCTAATGACT
GAACCCAGACATGGAACTCTTGAAGGATCCTTCTTACCACCCCAACCGAAACCTTGAACACGGGGATTGTGTAAATGACCGAGGAGATTACTGA 2400

pCB201 insert = U4

S V C C T F E N F L G R N G G V A F G N L C P L N T F T G L L . . L

TTGGGAAAAGATGATTCTGGGTCTTTCCCTTGACTTCTTGTTCATTACAACCTCTGGGCTTCTGGGGAGGGGTCAGAAAACATCAAAACACTGC
AACCCCTTTTCTACTAAGACCCAGAAAGGGAAC TGAAGAACAAGTTAATGTTTGAAGGACCCGAAAGACCCCTCCCAAGTCTTTGTAGTTTGTGACG 2500

pCB201 insert = U4

V G K D D S G S F P . L L V S I T N S V A F V G G V Q K T S K H C

AGCAGTTCCTAAATGATTCTCACAAGCAACCTGAGAGAGACAGTCTTGTGAGGGAGATCTGGGGGAGGCAGGAAGCTCCTCAGATTTTCTCACAGACCE
TCGTCAAGGATTTACTAAGAGTGTTCGTGGGACTCTCTCTGTCAGAACACTCCCTCTAGACCCCTCCGTCTTCGAGGAGTCTAAAAGAGTGTCTGGG 2600

pCB201 insert = U4

S S S . M I L T S N P E R D S L V R E I V G R O E A P O I F S Q T

TTCCCAATTCATCACCCTGCAACACTCGTCCGGAATCTGTCAGATATCCAGCACAGTGGCGGCCGCTCGAGTCTAGAGGGCCCGTTTAAACCCGCTG
AAGGGTTAAGGTAGTGGTGACGGTTGTGAGCAGGCTTAAGACGCTATAGGTCGTGTACCGCCGGCGAGCTCAGATCTCCCGGGCAAAATTTGGGCGAC 2700

pCB201 insert = U4

L P N S I T T A N T R P E F C R Y P A Q V R P L E S R G P V . T R .

ATCAGCTCGACTGTGCTTCTAGTTGCCAGCCATCTGTGTTTGCCCCCTCCCCGTGCTTCTTTGACCTGGAAGGTGCCACTCCCACTGTCTTTCC
TAGTCGGAGCTGACACGGAAGATCAACGGTCGGTAGACAACAACGGGGAGGGGGCACGGAAGGAAC TGGGACCTTCCACGGTGAGGGTGACAGGAAGG 2800

S A S T V P S S C O P S V V C P S P V P S L T L E G A T P T V L S

TAATAAATGAGGAAATTCATCGCATTGTCTGAGTAGGTGTCTATTCTTCTGGGGGTGGGGTGGGGCAGGACAGCAAGGGGGAGGATTGGGAAGACA
ATTATTTTACTCCTTTAACGTAGCGTAACAGACTCATCCACAGTAAGATAAGACCCCCACCCCAACCCGTCCTGTCTTCCCCCTCCTAACCTTCTGT 2900

. N E E I A S H C L S R C H S I L G G G V G Q D S K G E D V E D

ATAGCAGCATGCTGGGGATGCGGTGGGCTCTATGGCTTCTGAGGCGGAAAGAACAGCTGGGGCTCTAGGGGGTATCCCCACGCGCCCTTAGCGGGC
TATCGTCGCTACGACCCCTACGCCACCCGAGATACCAAGACTCCGCTTTCTTGGTCGACCCCGAGATCCCCCATAGGGGTGCGCGGGACATCGCCGCG 3000

N S R H A G D A V G S M A S E A E R T S V G S R G Y P H A P C S G A

ATTAAGCGCGGGGTGTGGTGGTTACGCGCAGCGTSACCGCTACACTTGCCAGCGCCCTAGCGCCCGCTCCTTTCTGCTTCTTCTCTTCTTCTCTGCG
TAATTCGCGCCGCCACACCACCAATGCGCGTCTGCTGCGATGTGAACGCTCGCGGGATCGCGGGCAGGAAAGCGAAAGAAAGGAAAGGAAAGAGCGG 3100

L S A A G V V V T R S V T A T L A S A L A P A P F A F F P S F L A

ACGTTCCGCGGCTTTCCCGTCAAGCTCTAAATCGGSCATCCCTTTAGGSTTCGATTTAGTGCTTTACGGCACCTCGACCCCAAAAACCTTGATTAGG
TCAAGCGCGCGAAAGGGGAGTTTGAGATTTAGCCCTTAGGGAAATCCCAAGGCTAAATCACGAAATGCCGTGGAGCTGGGGTTTTTGAACATACTC 3200

T F A G F P R O A L N R G I P L G F R F S A L R H L D P K K L D

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GTGATGGTTACGTTAGTGGGCCATCGCCCTGATAGACGGTTTTTCGCCCTTTGACGTTGGAGTCCACGTTCTTTAATAGTGGACTCTTGTTCCTCAACTGG
320
CACTACCAAGTGCATCACCCGGTAGCGGGACTATCTGCCAAAAAGCGGGAAGTCAACCTCAGGTGCAAGAAATTATCACCTGAGAACAAGTTTGGAC
G D G S R S G P S P . . T V F R P L T L E S T F F N S G L L F Q T G
AACAACACTCAACCTATCTCGGTCTATTCTTTGATTATAAGGGATTTGGGGATTTCCGCCCTATTGGTTAAAAATGAGCTGATTTAACAAAAATTT
340
TTGTTGTAGTTGGGATAGAGCCAGATAAGAAAACATAATATTCCTAAACCCCTAAAGCCGGATAACCAATTTTACTCGACTAAATTTGTTTTAA
T T L N P I S V Y S F D L . G I L G I S A Y W L K N E L I . Q K F
AACGCGAATTAATTCTGTGGAATGTGTGTCAGTTAGGTGTGGAAAGTCCCGAGGCTCCCGAGGCAGGCAGAAATGCAAGCATGCATCTCAATTAGT
360
TTGCGCTTAATTAAGACACCTTACACACAGTCAATCCACACCTTTTCAGGGGTCGAGGGGTCGCTCGTCTTCATACGTTTCGTACGTAGAGTTAATCA
N A N . F C G M C V S . G V E S P Q A P Q A G R S M Q S M H L N .
CAGCAACCGAGGTGGGAAAGTCCCGAGGCTCCCGAGCAGGCAGAAGTATGCAAGCATGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCC
380
GTCGTTGGTCCACACCTTTTCAGGGGTCGAGGGGTCGCTCGTCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTATCAGGCGGGGATTGAGG
S A T R C G K S P G S P A G R S M Q S M H L N . S A T I V P P L T P
GCCCATCCCGCCCTAACTCCGCCAGTTCGCCCATTCGCCCATGGCTGACTAATTTTTTTATTTATGCAGAGGCCGAGGCCGCTCTGCCTCT
370
CGGGTAGGGCGGGATTGAGGCGGGTCAAGCGGGTAAGAGCGGGGTACCGACTGATTAAAAAAATAAATACGCTCCGGCTCCGCGGAGACGGAGGA
P I P P L T P P S S A H S P P H G . L I F F I Y A E A E A A S A S
GAGCTATCCAGAAGTAGTGAGGAGGCTTTTTGGAGGCTTAGGCTTTTGCAAAAAGCTCCCGGGAGCTTGATATCCATTTTCGGATCTGATCAAGAGA
380
CTCGATAAGGTCTTCATCACTCTCCGAAAAACCTCCGGATCCGAAAAAGCTTTTCGAGGGCCCTCGAACATATAGGTAAAGCCCTAGACTAGTTCTCT
E L F Q K . . G G F F G G L G F C K K L P G A C I S I F G S D Q E
CAGATGAGGATCGTTTCGCATGATTGAACAAGAAGGATTCACGACGAGGTTCTCCGGCGCTTGGGTGGAGAGGCTATTCGGCTATGACTGGGCACAACA
390
GTCTTACTCTAGCAAGCGTACTAATCTGTTCTACCTAACGTGCGTCCAAGAGGCCGGCGAACCCACCTCTCCGATAAGCCGATCTGACCCGTGTTG
T S . G S F R M I E Q D G L H A G S P A A W V E R L F G Y D W A Q Q
GACAATCGGCTGCTCTGATGCCCGGTGTTCCGGCTGTCAGCGCAGGGGCGCCGGTCTTTTTTGCAAGACCGACCTGTCCGGTGCCTGAATGAAGT
400
CTGTTAGCGAGCAGACTACGGCGGCACAAGGCGACAGTCGGCTCCCGGGGCAAGAAAAACAGTTCTGGCTGGACAGGCCACGGGACTTACTTGAC
T I G C S D A A V F R L S A Q G R P V L F V K T D L S G A L N E L
CAGGACGAGGACGCGGGCTATCGTGGCTGGCCACGACGGGCGTTCCTTGGCAGCTGTGCTCGACGTTGTCTACTGAAGCGGGAAGGACTGGCTGCTA
410
GTCTGCTCCGTGCGCGGATAGCACCGACCGGCTGCTGCCGCAAGGAACGCGTGCACACGAGCTGCAACAGTACTTCGCCCCCTCCCTGACCGAGGATA
G D E A A R L S V L A T T G V P C A A V L D V V T E A G R D V L L
TGGCGAAGTGCCGCGGAGGATCTCTGTATCTCACCTTGTCTCTGCCGAGAAAGTATCCATCATGGCTGATGCAATGCGGCGGCTGCATAGCTTGG
420
ACCGGCTTCACGCGCCCGTCTAGAGGACAGTASAGTGAACGAGGACGGCTCTTCATAGGTAGTACCGACTACGTTACGCCCGGAGCTATGCGAAT
L G E V P S Q D L L S S H L A P A E K V S I M A D A M R R L H T L D
TCCGGCTACCTGCCCATTCGACCACCAAGCGAAACATCGCATCGAGCGAGCACGTACTCGGATGGAAGCCGGTCTTGTGATCAGGATGATCTGGACGAA
430
AGCCGATGAGGAGGCTAAGCTGGTGGTTTCGCTTTGAGCGTAGCTCGCTCGTGCATGAGGCTACCTTCGGCCAGAACAGCTAGTCTCTACTAGAGCTGCT
P A T C P F D H O A K H R I E R A R T R M E A G L V D Q D D L D E
GAGCATCAGGGCTCAGCGCCAGCGAAGTGTCTCCAGGCTCAAGGCGCGCATGCCGACGGCGAGGATCTCGTCTGACCCATGGCGATGCTGCTTGG
440
CTGCTAGTCCCGGAGCGGCTGGCTTGACAGCGGTCGAGTTCCGCGCTACGGGCTGCCGCTCCTAGAGCAGCACTGGGTACCGCTACGGAGCGAGG
E H G G L A P A E L F A R L K A R M P D G E D - L V V T H G D A C L

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CGAATATCATGGTGGAAAAATGGCCGCTTTTCTGGATTTCATCGACTGTGGCCGGCTGGGTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACCCGTGA
GCTTATAGTACCACCTTTTACCGGCGAAAAAGACCTAAGTAGCTGACACCGGCCGACCCACACCGCCTGGCGATAGTCCGTATCGCAACCGATGGGCACCT 2500
P N I M V E N G R F S G F I D C G R L G V A D R Y Q D I A L A T R D
TATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCGCTTCTCTGTGCTTTACGGTATCGCCGCTCCCGATTTCGAGCGCATCGCCTTCTATCGCCTTCTT
ATAACGACTTCTCGAACCGCCGCTTACCCGACTGGCGAAGGAGCACGAAATGCCATAGCGGCAGGGGCTAAGCGTCGCGTAGCGGAAGATAGCGGAAGAA 2600
I A E E L G G E V A D R F L V L Y G I A A P D S Q R I A F Y R L L
GACGAGTTCCTCTGAGCGGGACTCTGGGGTTCGAAATGACCGACCAAGCGACGCCCAACCTGCCATCACGAGATTCGATTCCACCGCCGCTTCTATGA
CTGCTCAAGAAGACTCGCCCTGAGACCCCAAGCTTTACTGGCTGGTTGCTGCGGGTGGACGGTAGTGTCTAAAGCTAAGGTGGCGGCGGAAGATACT 2700
D E F F . A G L V G S K . P T K R R P T C H H E I S I P P P P S M
AAGGTGGGCTTCGGAATCGTTTTCCGGGACGCCGGCTGGATGATCCTCCAGCGCGGGGATCTCATGCTGGAGTTCCTCGCCACCCCAACTTGTTTATT
TTCCAACCGAAGCCTTAGCAAAAGGCCCTGCGGCCGACCTACTAGGAGGTGCGGCCCTAGAGTACGACCTCAAGAAGCGGGTGGGTTGAACAAATAA 2800
K G V A S E S F S G T P A G . S S S A G I S C V S S S S P T P T C L L
GCAGCTTATAATGGTTACAAATAAGCAATAGCATCACAAATTTACAAATAAGCATTTTTTTCACTGCATTCTAGTTGTGGTTTGCCAAACTCATCA
CGTCGAATATTACCAATGTTTATTTTCGTTATCGTAGTGTTTAAAGTGTTTATTTTCGTAAGGAGTGAAGTCAACACCAACAGGTTTGAGTAGT 2900
Q L I M V T N K A I A S Q I S Q I K H F F H C I L V V V C P N S S
ATGTATCTTATCATGTCTGTATACCGTCGACCTCTAGCTAGAGCTTGGCGTAATCATGGTCATAGCTGTTTCTGTGTGAAATGTTATCCGCTCACAAT
TACATAGAATAGTACAGACATATGGCAGCTGGAGATCGATCTCGAACCGCATTAGTACCAGTATCGACAAAGGACACACTTTAACAATAGGCGAGTGTTA 3000
M Y L I M S V Y R R P L A R A V R N H G H S C F L C E I V I R S Q
TCCACACAACATACGAGCGGGAAGCATAAAGTGTAAGCCTGGGGTGCCATAGAGTGAGCTAACTCACATTAATTGCGTTG 5082
AGGTGTGTTGATGCTCGGCCCTTCGTATTTACATTTTCGGACCCACGGATTACTCACTCGATTGAGTGTAATTAACGCAAC
F H T T Y E P E A . S V K P G V P N E . A N S H . L R V

Claims

1. A vertebrate protein homologue of an UNC-53 protein of C. elegans or a functional equivalent, derivative or bioprecursor thereof, which protein comprises an amino acid sequence having a statistically significant homology to the amino acid sequence of said UNC-53 protein of C. elegans illustrated in Figure 2.
2. A vertebrate protein homologue of an UNC-53 protein of C. elegans, which protein comprises an amino acid sequence having one or more of sequence blocks A, B, C, D or E as illustrated in Figure 9a, or block F in Figure 12a or a sequence having a statistically significant homology therewith.
3. A vertebrate protein homologue of an UNC-53 protein of C. elegans, which protein comprises an amino acid sequence having one or more of sequence blocks A, B, C, D, E or F which differ from those blocks of Figure 9a or 12a only in conservative amino acid changes.
4. A vertebrate protein having an amino acid sequence encoded by the nucleotide sequence shown from nucleotide positions 1 to 6013 illustrated in Sequence ID No. 3.
5. A vertebrate protein comprising an amino acid sequence which comprises one or more of the prosite signatures as illustrated in Figure 28 for each of said sequences of homology as claimed in claim 2.
6. A vertebrate protein comprising an amino acid

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sequence as claimed in any one of claims 1 to 6 which is a human protein or a mouse protein.

5 7. A vertebrate protein having an amino acid sequence encoded by the nucleotide sequence shown in Sequence ID No. 4.

10 8. A vertebrate protein homologue according to any one of claims 1 to 7 comprising an amino acid sequence as shown in Sequence ID No. 1 or an amino acid sequence which differs from the amino acid sequence shown in Sequence ID No. 1 in one or more conservative amino acid changes.

15 9. A vertebrate protein homologue according to any one of claims 1 to 7 comprising an amino acid sequence as shown in Sequence ID No. 2 or an amino acid sequence which differs from the amino acid sequence shown in Sequence ID No. 2 in one or more
20 conservative amino acid changes.

25 10. A cDNA encoding a vertebrate homologue of UNC-53 protein of C. elegans according to any of claims 1 to 9.

30 11. A cDNA according to claim 10 comprising a sequence of nucleotides encoding an amino acid sequence as shown in Sequence ID No. 1 or an amino acid sequence which differs from the amino acid sequence shown in Sequence ID No. 1 only in one or more conservative amino acid changes.

35 12. A cDNA according to claim 10 comprising a sequence of nucleotides encoding an amino acid sequence as shown in Sequence ID No. 2 or an amino

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acid sequence which differs from the amino acid sequence shown in Sequence ID No. 2 only in one or more conservative amino acid changes.

5 13. A cDNA according to any of claims 10 or 11 which cDNA comprises at least from nucleotide position 1 to position 6013 of the sequence as shown in Sequence ID No. 3.

10 14. A cDNA according to claim 10 or 12 which comprises the nucleotide sequence illustrated in Sequence ID No. 4.

15 15. A nucleic acid molecule capable of hybridising to the DNA sequences according to any of claims 10 to 14 under high stringency conditions.

20 16. A DNA expression vector which comprises a cDNA as claimed any of claims 10 to 14.

25 17. A vector according to claim 16 which comprises a promoter of C. elegans UNC-53 protein or a vertebrate homologue thereof according to any of claims 1 to 9.

30 18. A vector according to claim 17 wherein said promoter sequence is derived from a gene encoding a mouse or human homologue of an UNC-53 protein of C. elegans.

35 19. A vector according to any of claims 16 to 18 which further comprises a sequence encoding a reporter molecule.

20. A vector according to claim 19 wherein said

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reporter molecule is a fluorophore.

21. A host cell transformed or transfected with the vector of any of claims 16 to 20.

5

22. A host cell transformed or transfected with the vector of claims 19 or 20.

23. A host cell according to claims 21 or 22,
10 which cell comprises a prokaryotic cell such as a bacterial cell or a eukaryotic cell such as a fungal, an animal, a plant or an insect cell.

24. A transgenic cell, tissue or organism
15 comprising a transgene capable of expressing a protein according to any of claims 1 to 9.

25. A transgenic cell, tissue or organism according to claim 24 which comprises any of a COS
20 cell, Hep G2, MCF-7 cell, N4 mouse neuroblastoma cell, a NIH3T3 cell, or colorectal carcinoma or human derived cells.

26. A transgenic cell, tissue or organism
25 according to claim 24 or 25 wherein said transgene comprises a vector according to any of claims 16 to 20.

27. A transgenic cell, tissue or organism
30 according to claim 24 to 26 wherein said transgene comprises a vector according to claim 19 or 20.

28. A transgenic cell, tissue or organism according to any of claims 24 to 26 wherein said
35 organism comprises any of an insect, a fungus, a non-

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human mammal, a plant or a nematode worm.

29. A method of producing a mutant vertebrate non-human organism which mutation affects cell
5 behaviour or the regulation of cell motility or the shape or the direction of cell migration, which method comprises inducing a mutation in the wild type gene encoding the vertebrate homologue of an UNC-53
10 C. elegans protein.

30. A vertebrate protein homologue of an UNC-53 protein of C. elegans, or a functional equivalent, derivative, fragment or bioprecursor thereof, for use
15 as a medicament to promote neuronal regeneration, revascularisation, wound healing or for treatment of chronic neuro-degenerative diseases or acute traumatic injuries or fibrotic disease.

31. A vertebrate protein homologue of an UNC-53
20 protein of C. elegans for use as claimed in claim 30 wherein said vertebrate human homologue is as claimed in any one of claims 1 to 9.

32. Use of a vertebrate protein homologue of an
25 UNC-53 protein of C. elegans, or a functional equivalent, derivative, fragment or bioprecursor thereof, in the manufacture of a medicament for promoting neuronal regeneration, revascularisation, wound healing or for treatment of chronic
30 neurodegenerative diseases or acute traumatic injuries or fibrotic disease.

33. Use of a vertebrate protein homologue of
35 UNC-53 protein of C. elegans according to claim 32 wherein said vertebrate protein homologue is as

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claimed in any one of claims 1 to 9.

34. A pharmaceutical composition comprising a vertebrate homologue of an UNC-53 protein of C. elegans, or a functional equivalent, derivative, fragment or bioprecursor of said vertebrate protein, together with a pharmaceutically acceptable carrier, diluent or excipient therefor.

35. A pharmaceutical composition as claimed in claim 34 which comprises a vertebrate homologue of an UNC-53 protein of C. elegans according to any of claims 1 to 9.

36. A nucleic acid sequence encoding a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, fragment, derivative or bioprecursor of said vertebrate homologue, for use as a medicament.

37. A nucleic acid sequence according to claim 36 wherein said sequence is a cDNA sequence as claimed in any of claims 10 to 14 or a functional fragment of said cDNA sequence.

38. Use of a nucleic acid sequence encoding a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, fragment, derivative or bioprecursor of said vertebrate homologue, in the manufacture of a medicament to promote neuronal regeneration, revascularisation or wound healing, or for treatment of chronic neurodegenerative diseases or acute traumatic injuries or fibrotic disease.

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39. Use of a nucleic acid sequence according to claim 38 wherein said sequence is a cDNA sequence as claimed in any of claims 10 to 14 or a functional fragment of said nucleic acid sequence.

5

40. A pharmaceutical composition comprising a nucleic acid sequence according to claim 36 or 37 and a pharmaceutically acceptable carrier, diluent or excipient therefor.

10

41. A pharmaceutical composition according to claim 40 wherein said nucleic acid sequence is a cDNA sequence as claimed in any of claims 10 to 14.

15

42. A method of determining whether a compound is an inhibitor or enhancer of the regulation of cell behaviour, growth, cell shape or motility or the direction of cell migration, which method comprises contacting said compound with a host cell according to claim 21 or 23 or a transgenic cell as claimed in any of claims 24 to 27 and screening for a phenotypic change in said cell.

20

43. A method according to claim 41 which is capable of determining whether said compound is an inhibitor or an enhancer of the signal transduction pathway of said transgenic cell of which said vertebrate homologue of an UNC-53 protein or a functional equivalent, derivative, fragment or bioprecursor of said vertebrate homologue is a component or is an inhibitor or an enhancer of a parallel or redundant signal transduction pathway in said cell.

30

44. A method according to claim 43 wherein said

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method is capable of determining whether said compound is an inhibitor or an enhancer of said vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, fragment, derivative or bioprecursor of said vertebrate homologue.

45. A method according to any of claims 42 to 44 wherein said phenotypic change to be screened is a change in cell growth, or shape or a change in cell motility.

46. A method according to any of claims 42 to 44 wherein said phenotypic change to be screened is a change in filopodia outgrowth, ruffling behaviour, cell adhesion, contact inhibition or the length of neurite growth.

47. A method as claimed in any of claims 42 to 44 wherein said transgenic cell is an N4 neuroblastoma cell and the phenotypic change to be screened is the length of neurite growth.

48. A method as claimed in any of claims 42 to 44 wherein said transgenic cell is an MCF-7 breast carcinoma cell or an NIH3T3 cell and the phenotypic change to be screened is the extent of phagokinesis or contact inhibition.

49. A method of determining whether a compound is an inhibitor or an enhancer of the regulation of cell shape, cell growth or motility or of the direction of cell migration, which method comprises administering said compound to a transgenic organism according to any of claims 24 to 28 or a mutant organism produced according to the method of claim 29

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and screening for a phenotypic change in said organism.

50. A method according to claim 49, wherein said
5 method is capable of determining whether said compound
is an inhibitor or enhancer of a protein of the signal
transduction pathway of said transgenic or mutant
organisms, of which the vertebrate homologue of UNC-53
protein of C. elegans or a functional equivalent,
10 derivative, fragment or bioprecursor of said
vertebrate homologue is a component, or is an
inhibitor or an enhancer of a parallel or redundant
signal transduction pathway in said cell.

51. A method according to claim 50 wherein said
15 method is capable of determining whether said compound
is an inhibitor or an enhancer of the vertebrate
homologue of UNC-53 protein itself or a functional
equivalent, fragment, derivative or bioprecursor of
20 said vertebrate homologue.

52. A compound which is identifiable by the
method according to any one of the claims 42 to 51 as
an enhancer of the regulation of cell shape, or growth
25 or motility or the direction of cell migration for use
as a medicament for promoting neuronal regeneration,
revascularisation or wound healing or for treatment of
chronic neurodegenerative diseases or acute traumatic
injuries or fibrotic disease.

53. Use of a compound which is identifiable by
30 the method according to any one of the claims 42 to 51
as an enhancer of the regulation of cell shape, or
growth or motility or the direction of cell migration
35 in the preparation of medicament for promoting

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neuronal regeneration, revascularisation or wound healing or for treatment of chronic neurodegenerative diseases or acute traumatic injuries or fibrotic disease.

5

54. A pharmaceutical composition comprising a compound identified according to the method of any of claims 42 to 51 claim and a pharmaceutically acceptable carrier, diluent or excipient therefor.

10

55. A compound which is identifiable by the method according to any one of claims 42 to 51 as an inhibitor of the regulation of cell motility, growth, or shape, or the direction of cell migration, for use as a medicament for alleviating the spread of disease inducing cells or metastasis or loss of contact inhibition.

15

56. Use of a compound according to claim 55 in the manufacture of a medicament for alleviating the spread of disease inducing cells or metastasis or loss of contact inhibition.

20

57. A pharmaceutical composition comprising the compound as claimed in claim 55, and a pharmaceutically acceptable carrier diluent or excipient therefor.

25

58. A method of determining whether a compound is an inhibitor or an enhancer of transcription of a gene encoding a vertebrate homologue of UNC-53 protein of C. elegans, which method comprises the steps of (a) contacting said compound with a cell according to any of claims 22 or 27 and (b) monitoring the level of said reporter molecule and comparing the results

30

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obtained from said monitoring step with a control comprising a cell according to claims 22 or 27, which cell has not been contacted with said compound.

5 59. A method as claimed in claim 58 wherein said reporter molecule detected is mRNA or green fluorescent protein.

10 60. A compound which is identifiable by the method according to claims 58 or 59, as an enhancer of transcription of a gene coding for a vertebrate homologue of an UNC-53 protein of C. elegans or a functional fragment of said gene, for use in promoting neuronal regeneration, revascularisation or wound
15 healing, or for treatment of chronic neuro-degenerative diseases or acute traumatic injuries or fibrotic disease.

20 61. Use of a compound which is identifiable by the method of claims 58 or 59, as an enhancer of transcription of a gene coding for a vertebrate homologue of an UNC-53 protein of C. elegans or a functional fragment of said gene, in the manufacture of a medicament for promoting neuronal regeneration,
25 revascularisation or wound healing, or for treatment of chronic neuro-degenerative diseases or acute traumatic injuries or fibrotic disease.

30 62. A pharmaceutical composition which comprises the compound of claim 60 and a pharmaceutically acceptable carrier, diluent or excipient therefor.

35 63. A compound which is identifiable by the method of claims 58 or 59 as an inhibitor of transcription of a gene coding for a vertebrate

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homologue of a UNC-53 protein of C. elegans or a functional fragment of said gene for use in alleviating the spread of disease inducing cells or metastasis or loss of contact inhibition.

5

64. Use of a compound which is identifiable by the method of claims 58 or 59 as an inhibitor of transcription of a gene coding for a vertebrate homologue of an UNC-53 protein of C. elegans or a functional fragment of said gene, in the manufacture of a medicament for alleviating spread of disease inducing cells or metastasis or loss of contact inhibition.

65. A pharmaceutical composition which comprises the compound of claim 63 and a pharmaceutically acceptable carrier, diluent or excipient therefor.

66. A kit for determining whether a compound is an enhancer or an inhibitor of the regulation of cell motility, growth or shape or the direction of cell migration which kit comprises at least one transgenic cell as claimed in any one of claims 22 to 25 to be contacted with said compound and at least one cell according to claims 21 to 28 to be used as a control and means for contacting said compound with one of said at least one transgenic cells.

67. A kit for determining whether a compound is an inhibitor or an enhancer of transcription of a gene coding for a vertebrate homologue of an UNC-53 protein of C. elegans or a functional fragment of said gene which kit comprises at least one cell as claimed in any one of claims 21 to 25 means for contacting said compound with said cells.

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68. A kit for determining whether a compound is an enhancer or an inhibitor of the activity of a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, derivative, fragment or bioprecursor of said vertebrate homologue protein, which kit comprises at least, one vertebrate mutant non-human organism produced according to the method as claimed in claim 29 or a transgenic organism as claimed in claims 24 to 28 and a wild type of said vertebrate mutant organism.

69. A method of identifying vertebrate homologues of an unc-53 gene of C. elegans or a functional fragment thereof, which method comprises hybridizing to a DNA library a suitable oligonucleotide sequence of between 15 to 50 nucleotides of the nucleic acid sequence encoding unc-53 or a functional equivalent, derivative or bioprecursor thereof, under appropriate conditions of stringency to identify genes having statistically significant homology with the cDNA according to any of claims 10 to 14.

70. A method of identifying a protein which is active in the signal transduction pathway of a cell of which a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, fragment or bioprecursor of said vertebrate homologue is a component, which method comprises:

- (a) contacting an extract of said cell with an antibody to the vertebrate homologue of the UNC-53 protein of C. elegans or a functional equivalent, fragment, derivative or bioprecursor of said protein,
- (b) identifying the antibody/vertebrate

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homologue complex, and

(c) analysing the complex to identify any protein bound to the vertebrate homologue of UNC-53 protein of C. elegans other than the antibody.

5

71. A method of identifying a further protein which is active in the signal transduction pathway of a cell of which a vertebrate homologue of an UNC-53 protein or a functional equivalent, fragment or bioprecursor of said UNC-53 protein is a component, which method comprises:

10

(a) forming an antibody to the first identified protein bound to the vertebrate homologue of UNC-53 protein of C. elegans in claim 70,

15

(b) contacting a cell extract with said antibody and identifying the antibody/protein complex,

20

(c) analysing the complex to identify any further protein bound to the first protein other than the antibody, and

(d) optionally repeating steps (a) to (c) to identify further proteins in said pathway.

25

72. A method of identifying a protein which is active in the signal transduction pathway of a cell of which a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, fragment or bioprecursor of said homologue protein is a component, which method comprises

30

(a) contacting an extract of said cell with the vertebrate homologue of an UNC-53 protein of C. elegans or a functional

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equivalent, derivative or bioprecursor of said vertebrate homologue,

(b) identifying any vertebrate homologue of UNC-53 protein/protein complex formed and

5 (c) analysing the complex to identify any protein bound to the vertebrate homologue of UNC-53 protein other than the same vertebrate homologue of UNC-53 protein.

10 73. A method according to claim 72 which further comprises contacting a cell extract with any protein identified from step (c) not being the same as the vertebrate homologue of UNC-53 protein used and repeating steps (b) and (c) so as to identify any
15 further protein involved in the signal transduction pathway of said cell.

20 74. A method of identifying a protein involved in the signal transduction pathway of a cell of which a vertebrate homologue of an UNC-53 protein of C. elegans is a component which method comprises:

(a) providing an appropriate host cell having a DNA construct comprising a reporter gene under the control of a promoter
25 regulated by a transcription factor having a DNA binding domain and an activating domain,
(b) expressing in said host cell a first hybrid DNA sequence encoding a first fusion
30 of a fragment or all of a DNA sequence according to any of claims 10 to 14 and either said DNA binding domain or the activating domain of the transcription factor,
(c) expressing in the host cell at least
35 one second hybrid DNA sequence encoding a

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putative binding protein to be investigated together with the DNA binding or activating domain of the transcription factor which is not incorporated in the first fusion,

5 (d) detecting any binding of the protein being investigated with a protein according to any of claims 1 to 9 by detecting for the production of any reporter gene product in said host cell.

10

75. A protein identified by the method of any one of claims 70 to 74 for use as a medicament to promote neuronal regeneration, revascularisation or wound healing, or for treatment of chronic neuro-
15 degenerative diseases or acute traumatic injuries or fibrotic disease.

76. Use of a protein identified by the methods of any one of claims 70 to 74 in the manufacture of a
20 medicament for promoting neuronal regeneration, revascularisation or wound healing, or for treatment of chronic neurodegenerative diseases or acute traumatic injuries or fibrotic disease.

25 77. A pharmaceutical composition comprising a protein identified by the methods of any one of claims 70 to 74 and a pharmaceutically acceptable carrier, diluent, or excipient therefor.

30 78. A process for producing a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent fragment, derivative or bioprecursor of said vertebrate homologue which process comprises culturing the cells of any of claims
35 21 to 28 and recovering said vertebrate homologue of

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UNC-53 protein expressed.

79. A process for producing a vertebrate
homologue of an UNC-53 protein of C. elegans or a
5 functional equivalent, fragment, derivative or
bioprecursor of said protein which process comprises
culturing an insect cell transfected with a
recombinant Baculovirus vector, said vector comprising
a DNA insert encoding said vertebrate homologue of
10 UNC-53 protein or a functional equivalent, fragment or
bioprecursor of said vertebrate homologue, downstream
of the Baculovirus polyhedrin promoter, and recovering
the expressed vertebrate homologue of UNC-53 protein.

15 80. A nucleotide sequence comprising the
sequence as shown in figure 15.

81. A nucleotide sequence comprising the
sequence as shown in figure 16.

20 82. A nucleotide sequence comprising the
sequence as shown in figure 17.

83. A method of detecting whether a compound is
25 an inhibitor or an enhancer of expression of a
vertebrate homologue of an UNC-53 of C. elegans, or a
functional equivalent, derivative or fragment of said
vertebrate homologue which method comprises contacting
a cell expressing said homologue with said compound
30 and monitoring for a phenotypic change compared to a
control cell which has not been contacted with said
compound.

84. A method according to claim 83 wherein said
35 cell comprises a cell according to any of claims 21 to

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28.

85. A method according to claim 83 wherein said cell has undergone loss of contact inhibition.

5

86. A method according to any of claims 83 to 85 which is capable of determining whether said compound is an inhibitor of expression of said vertebrate homologue in which the compound to be tested comprises a nucleic acid.

10

87. A method according to claim 86 wherein said nucleic acid sequence comprises an antisense DNA or RNA sequence.

15

88. A method according to claim 87 wherein said mRNA sequence comprises 3' untranslated regions of mRNA encoding for said vertebrate homologue.

20

89. A method according to any of claims 83 to 85 wherein said compound to be tested comprises a protein having an amino acid sequence potentially suitable for inhibiting function of said vertebrate homologue.

25

90. A method according to claim 89 wherein said protein comprises a protein identified according to any of the methods of claims 70 to 74.

30

91. A pharmaceutical composition comprising a compound identified according to any of claims 83 to 89 together with a pharmaceutically acceptable carrier, diluent or excipient therefor.

35

92. A nucleic acid sequence identified according to the method of any of claims 86 to 88 for use in

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treatment of loss of contact inhibition or carcinoma which is mediated by a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, fragment, derivative or bioprecursor thereof.

93. Use of a nucleotide sequence identified according to the method of any one of claims 86 to 88 in the preparation of a medicament for the treatment of loss of contact inhibition or carcinoma which is mediated by a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, fragment, derivative or bioprecursor of said vertebrate homologue.

94. A nucleic acid according to claim 92 for use in the preparation of a medicament for inhibiting expression of a gene coding for a vertebrate homologue of an UNC-53 protein of C. elegans.

95. A NIH3T3 cell line transfected with pcB201 and deposited under LMBP Accession No. 1603CB.

96. A plasmid pcB 201 of Sequence ID No. 10 deposited under LMBP Accession No. LMBP 3594.

97. A MCF-7 cell line transfected with plasmid pcB 201 deposited under LMBP Accession No. LMBP 1601CB.

98. An assay for detecting expression of a vertebrate homologue of UNC-53 protein of C. elegans in a vertebrate cell which assay comprises contacting a cell or an extract thereof with an antibody to said vertebrate homologue, or a functional equivalent,

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derivative or bioprecursor thereof, which antibody is linked to a reporter molecule, removing any unbound antibody and monitoring for the presence of said reporter molecule.

5

99. An assay according to claim 98 wherein said reporter molecule is an antibody conjugated with a suitable fluorophore or detectable enzyme.

10

100. A method for detecting for expression of a gene coding for a vertebrate homologue of an UNC-53 protein of C. elegans or a functional equivalent, derivative, fragment or bioprecursor thereof, which method comprises contacting a probe specific for a nucleic acid or protein sequence coding for or corresponding to said vertebrate homologue or a functional equivalent, fragment or bioprecursor therefor with a cell extract which probe is linked to a reporter and analysing for the presence of said reporter.

20

101. A method according to claim 100 wherein said probe comprises a complimentary sequence to a region of mRNA transcribed from said gene encoding said vertebrate homologue of UNC-53 protein or a functional equivalent, derivative or bioprecursor therefor.

25

102. A method according to claim 101 wherein said complimentary sequence is a 3' or 5' untranslated region of said mRNA.

30

103. A method according to claims 100 or 102 wherein said reporter comprises a radiolabel.

35

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104. A method according to claim 100 wherein
said probe comprises an antibody specific for said
vertebrate homologue of said UNC-53 protein or a
functional equivalent, derivative, fragment or
5 bioprecursor therefor.

105. A method according to claim 104 wherein
said reporter comprises an antibody conjugated with a
detectable fluorophore or enzyme.
10

106. Phage Lambda clone 3b of Sequence ID No. 5
deposited under Accession No. LMBP 3595.

107. A method of determining whether a compound
15 is an inhibitor or an enhancer of association of UNC-
53 or a vertebrate homologue thereof according to any
of claims to 1 to 9 to microtubules or plus end
regions thereof, which method comprises:-

(a) contacting said compound with a
20 transgenic cell, tissue or organism
expressing UNC-53 protein or said vertebrate
homologue and which protein is operably
linked to a reporter molecule.

(b) screening for the localisation of said
25 reporter molecule as compared to a cell
according to step (a) which has not been
contacted with said compound.

108. A compound identifiable by the method
30 according to claim 107.

109. A compound identifiable by the method
according to claim 107 as an inhibitor of localisation
or association of UNC-53 or said vertebrate homologue
35 with microtubules or the plus end region thereof for

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use in alleviating the spread of disease inducing cells or metastasis or loss of contact inhibition.

110. A compound identifiable by the method
5 according to claim 107 as an enhancer of association of UNC-53 or said vertebrate homologue with microtubules or the plus end region thereof, for use in promoting neuronal regeneration, revascularisation or wound healing, or for treating chronic
10 neurodegenerative diseases or acute traumatic injuries or fibrotic disease.

111. A pharmaceutical composition comprising the compound according to claims 108 or 109 and a
15 pharmaceutically acceptable carrier, diluent or excipient therefor.

112. A kit for determining whether a compound is an inhibitor or an enhancer of association of UNC-53
20 or a vertebrate homologue thereof according to any of claims 1 to 9 with microtubules or the plus end regions thereof, which kit comprises at least one transgenic cell expressing UNC-53 and a reporter molecule or a cell according to any of claims 20 to 24
25 and at least one cell of the same cell type for use as a control and means for contacting said compound with one of said at least one transgenic cells.

113. A composition comprising UNC-53 of C. elegans
30 elegans or a vertebrate homologue thereof according to any of claims 1 to 9 linked to a compound identified as an inhibitor or enhancer of association of UNC-53 or said vertebrate homologue with microtubules or their plus end regions for use in targeting said
35 compound to said microtubule or the plus end regions

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thereof.

114. A composition according to claim 113 which further comprises a cell transformation or
5 transfecting agent.

115. A method of targeting a protein to a cell microtubule or the plus end region thereof, which method comprises introducing into a host cell, tissue
10 or organism a transgene comprising a sequence capable of expressing UNC-53 or a vertebrate homologue thereof according to any of claims 1 to 9, which sequence is operably linked to a sequence encoding said protein to be targeted such that a chimeric protein is expressed
15 and which results in targeting said protein to said microtubule or a plus end region thereof.

116. A method of identifying a molecule which covalently modifies UNC-53 or a vertebrate homologue
20 thereof according to any of claims 1 to 9, which method comprises

a) contacting either an extract from a cell expressing UNC-53 or said vertebrate homologue or a mixture of enzymes comprising candidate UNC-53
25 modifying enzymes in the presence of an indicator of covalent modification of a protein,

b) identifying any covalently modified UNC-53 protein from step a),

c) identifying said molecule involved in said
30 modification step.

117. A method according to claim 112, wherein said indicator comprises ¹⁴C-p.

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118. A method of identifying a compound which alleviates or enhances the toxicity of UNC-53 or a vertebrate homologue thereof according to any of claims 1 to 9, which method comprises contacting said compound with a cell, tissue or organism according to claim 27, and monitoring for the presence of said reporter molecule adjacent said microtubules or the plus end regions thereof.

119. Plasmid pLM1 of Sequence ID No. 6 deposited under Accession No. LMBP 3762.

120. Plasmid pLM4 of Sequence ID No. 7 deposited under Accession No. LMBP 3763.

121. Plasmid pEGF72 of Sequence ID No. 8 deposited under Accession No. LMBP 3764.

122. Plasmid pCB501 of Sequence ID No. 9 deposited under LMBP Accession No. LMBP 3765.

123. A worm strain comprising a chimeric C.elegans human unc-53 gene deposited under LMBP Accession No. LMBP-1663CB.

124. A vertebrate homologue according to any of claims 1 to 3 which is a mouse homologue.

125. A homologue according to claim 125 having the sequence illustrated in Figure 14.

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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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AlwI

ACCTCGCTCTGCTAATCCTGTTACCAGTGGCTGCTGCCAGTGGCGATAAGTCGTGCTTACCGGGTTGGACTCAAGACGATAGTTACCGGATAAGGCGCA 9697
TGGAGCGAGACGATTAGGACAATGGTCACCGACGACGGTCACCGCTATTACGACACAGAATGGCCCAACCTGAGTTCTGCTATCAATGGCCTATTCCGCGT
P R S A N P V T S G C C Q V R . V V S Y R V G L K T I V T G . G A

Apal I

GCGGTCGGGCTGAACGGGGGGTTCGTGCACACAGCCAGCTTGGAGCGAACGACCTACACGAACTGAGATACCTACAGCGTGAGCTATGAGAAAGCGCC 9700
CGCCAGCCCGACTTGCCCCCAAGCAGTGTGTGCGGTCGAACCTCGCTTGTGATGTGGCTTGACTCTATGGATGTGCGACTCGATACTCTTTCGCGG
A V G L N G G F V H T A Q L G A N D L H R T E I P T A . A M R K R

ACGCTTCCCGAAGGGAGAAAGCGGACAGGTATCCGGTAAGCGGCAGGGTCGGAACAGGAGAGCGCACGAGGGAGCTTCCAGGGGGAAACGCCTGGTATC 9703
TGCGAAGGGCTTCCCTCTTTCGCGCTGTCCATAGGCCATTGCGCGTCCCAGCCTTGTCTCTCGCGTGCCTCCCGAAGGTCCCCCTTTGCGGACCATAA
H A S R R E K G G Q V S G K R Q G R N R R A H E G A S R G K R L V S

TTTATAGTCCTGTCGGGTTTCGCCACCTCTGACTTGAGCGTCGATTTTGTGATGCTCGTCAGGGGGGCGGAGCCTATGGAAAAACGCCAGCAACGCGGC 9706
AAATATCAGGACAGCCCAAAGCGGTGGAGACTGAACTCGCAGCTAAAAACACTACGAGCAGTCCCCCGCCTCGGATACCTTTTTCGCGTCTGTCGCGC
L . S C R V S P P L T . A S I F V M L V R G A E P M E K R Q Q R G

Ava III
Nsi I

CTTTTACGGTTCTTGGCCTTTTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 9709
GAAAAATGCCAAGGACCGGAAACGACCGGAAACGAGTGTAACAAGAAAGGACGCAATAGGGGACTAAGACACCTATTGGCATAATGGCGGTACGTA
L F T V P G L L L A F C S H V L S C V I P . F C G . P Y Y R H A

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fig 31 pEGFPsma (1 > 6960) Site and Sequence

Enzymes: 72 of 146 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

Page 1

TAGTTATTAATAGTAATCAATTACGGGGTCATTAGTTTCATAGCCCATATATGGAGTTCGCGTTACATAAATTACSGTAAATGGCCCGCTGGCTGACCG
ATCAATAATTATCATTAGTTAATGCCCCAGTAATCAAGTATCGGGTATATACCTCAAGGCGCAATGTATGAATGCCATTACCGGGCGGACCGACTGGC
L L I V I N Y G V I S S . P I Y G V P R Y I T Y G K V P A V L T
CCCAACGACCCCGCCCATTGACGTCAATAATGACGTATGTTCCCATAGTAACGCCAATAGGGACTTTCATTGACGTCAATGGGTGGAGTATTTACGGT
GGGTTGCTGGGGGCGGGTAACCTGCAGTTATTACTGCATACAAGGGTATCATTCGCGTTATCCCTGAAAGGTAACCTGCAGTTACCCACCTCATAAATGCCA
A Q R P P P I D V N N D V C S H S N A N R D F P L T S M G G V F T V
AAACTGCCCACTTGGCAGTACATCAAGTGTATCATATGCCAAGTACGCCCTTATGACGTCAATGACGGTAAATGGCCCGCTGGCATTATGCCAGTA
TTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTCATCGGGGGGATAACTGCAGTTACTGCCATTACCGGGCGGACCGTAATACGGGTCA
N C P L G S T S S V S Y A K Y A P Y . R Q . R . M A R L A L C P V
CATGACCTTATGGGACTTTCTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGATGCGGTTTTGGCAGTACATCAATGGCGTGGG
GTACTGGAATACCTGAAAGGATGAACCGTCATGTAGATGCATAATCAGTAGCGATAATGGTACCACCTACGCCAAAACCGTCATGTAGTTACCCGACCT
H D L M G L S Y L A V H L R I S H R Y Y H G D A V L A V H O V A V
TAGCGGTTTGACTCACGGGGATTCCAAGTCTCCACCCATTGACGTCAATGGGAGTTTGTGTTTGGCACAAAATCAACGGGACTTTCCAAAATGTCGTA
ATCGCCAAACTGAGTGCCCTAAAGGTTAGAGGTGGGGTAACCTGCAGTTACCTCAAAACAAACCGTGGTTTTAGTTGCCCTGAAAGGTTTTACAGCA
I A V . L T G I S K S P P H . R Q V E F V L A P K S T G L S K M S
ACAACCTCGCCCATTGACGCAATGGGCGGTAGGCGGTACGGTGGGAGGTCATATAAGCAGAGCTGGTTTTAGTGAACCGTCAGATCCGCTAGCGCTA
TGTTGAGGCGGGGTAACCTGCGTTTACCGCCATCCGCACATGCCACCTCCAGATATATTCGCTCGACCAAACTACTGGCAGCTAGGCGATCGCGA
O L R P I D A N G R . A C T V G G L Y K Q S V F S E P S D P L A L
CCGGTCGCCACCATGGTGAGCAAGGGCGAGGAGCTGTTACCGGGGTGGTGCCCATCCTGGTCGAGCTGGACGGCGACGTAACGGCCACAAGTTCAGCG
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eGFPc.e.unc53sma
P V A T M V S K G E E L F T G V V P I L V E L D G D V N G H K F S
TGTCGGCGAGGGCGAGGGCGATGCCACCTACGGCAAGCTGACCTGAAGTTCATCTGCACCACCGGCAAGCTGCCCGTGCCCTGGCCCAACCTCGTGAC
ACAGGCGCTCCCGCTCCCGCTACGGTGGATGCCGTTGACCTGGGACTTCAAGTAGACGTGGTGGCGGTCGACGGGCACGGGACCGGTGGGAGCACTG
eGFPc.e.unc53sma
V S G E G E G D A T Y G K L T L K F I C T T G K L P V P V P T L V T
CACCTGACCTACGGCGTGCAGTGCTTCAGCGCTACCCCGACCACATGAAGCAGCAGCACTTCTCAAGTCCGCCATGCCCGAAGGCTACGTCCAGGAG
GTGGGACTGGATGCCGACGTCACGAAGTCGGCGATGGGGCTGGTGTACTTCGTCGTCTGAAGAAGTTCAGGCGGTACGGGCTTCGGATGCAGGTCCTC
eGFPc.e.unc53sma
T L T Y G V Q C F S R Y P D H M K Q H D F F K S A M P E G Y V Q E
CGCACCATCTTCTTCAAGGACGACGGCAACTACAAGACCGCGCCGAGGTGAAGTTCGAGGGCGACACCTGGTGAAACCGCATCGAGCTGAAGGSCATCG
GCGTGGTAGAAGAAGTTCCTGCTGCCGTTGATGTTCTGGGCGCGGC TCCACTTCAAGCTCCCGCTGTGGGACCACTTGGCGTAGCTCGAATTCCCGTAGC
eGFPc.e.unc53sma
P T I F F K D D G N Y K T R A E V K F E G D I L V H R I E L K G I

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fig 31 pEGFPsma (1 > 6960) Site and Sequence

Page 2

ACTTCAAGGAGGACGGCAACATCCTGGGGCACAAGCTGGAGTACAAC TACAAGCCACAACGCTATATCATGGCCGACAAGCAGAAGAACGGCATCAA
TGAAGTTCTCTCCGCGTTGTAGGACCCCGTGTTCGACCTCATGTTGATGTTGTCGGTGTTCAGATATAGTACCGGCTGTTCTGCTCTCTTGGCGTAGTT 1100

eGFPc.e.unc53sma

D F K E D G N I L G H K L E Y N Y N S H N V Y I M A D K O K N G I I

GGTGAACITCAAGATCCGCCACAACATCGAGGACGGCAGCGTGCAGCTCGCCGACCACTACCAGCAGAACACCCCATCGGCACGCGCCCGTGTCTGCTG
CCACTTGAAGTTCTAGGCGGTGTGTAGCTCTGCCGTCGCACGTCGAGCGGCTGGTGTGGTCTGTTGTGGGGGTAGCCGCTGCCGGGGCACGACGAC 1200

eGFPc.e.unc53sma

V N F K I R H N I E D G S V Q L A D H Y Q Q N T P I G D G P V L L

CCCGACAACCAC TACCTGAGCACCCAGTCCGCCCTGAGCAAGACCCCAACGAGAAGCGCGATCACATGGTCTCTGCTGGAGTTCGTGACCGCCGCCGGGA
GGGCTGTTGGTGTATGGACTCGTGGGTCAGGCGGGACTCGTTTCTGGGGTTGCTCTTCGCGCTAGTGTACCAGGACGACCTCAAGCACTGGCGGGCGGCCCT 1300

eGFPc.e.unc53sma

P D N H Y L S T O S A L S K D P N E K R D H M V L L E F V T A A G

TCACTCTCGGCATGGACGAGCTGTACAAGTCCGGACTCAGATCTACGTCAAATGTAGAATTGATACCAATCTACACGGATTGGGCCAATCGGCACCTTTT
AGTGAGAGCGGTACCTGCTCGACATGTTTCAGGCCTGAGTCTAGATGCAGTTTACATCTTAACATATGGTTAGATGTGCTTAACCCGGTTAGCCGTGGAAAG 1400

eGFPc.e.unc53sma

C.e.unc53 sma

I T L G M D E L Y K S G L R S T S N V E L I P I Y T D W A N R H L S

GAAGGGCAGCTTATCAAGTCGATTAGGGATATTTCCAATGATTTTCGCGACTATCGACTGGTTTCTCAGCTTATTAATGTGATCGTTCCGATCAACGAA
CTTCCCGTCGAATAGTTTCAGCTAATCCCTATAAAGGTTACTAAAAGCGCTGATAGCTGACCAAGAGTCGAATAATTACACTAGCAAGGCTAGTTGCTT 1500

eGFPc.e.unc53sma

C.e.unc53 sma

K G S L S K S I R D I S N D F R D Y R L V S Q L I N V I V P I H E

TTCGCGCTGCATTACGAAACGTTTGGCAAAAATCACATCGAACC TGGATGGCTCGAAACGTGTC TCGACTACCTGAAAAATCTGGGCTCTGACTGCT
AAGAGCGGACGTAAGTGCTTTGCAAACCGTTTGTAGTGTAGCTTGGACCTACCGGAGCTTTCACAGAGCTGATGGACTTTTGTAGCCAGAGCTGACGA 1600

eGFPc.e.unc53sma

C.e.unc53 sma

F S P A F T K R L A K I T S N L D G L E T C L D Y L K N L G L D C

CGAAACTCACAAAACCGATATCGACAGCGGAAACTTGGGTGCAGTTC TCCAGCTGCTCTTCCTGCTCTCCACCTACAAGCAGAAGCTTCGGCAACTGAA
GCTTTGAGTGGTTTTGGCTATAGCTGTCGCCCTTTGAACCCACGTCAAGAGGTCGACGAGAAGGACGAGAGGTGGATGTTCTGCTTTCGAAGCCGTTGACTT 1700

eGFPc.e.unc53sma

C.e.unc53 sma

S X L T K T D I D S S Y L G A V L Q L L F L L S T Y K Q K L R Q L I

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fig 31 pEGFPsma (1 > 6960) Site and Sequence

Page 1

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TTTTCTAGTCTTCTTTAACCTCGTTGATGGGTGATAGGTAATACGGTGGGCGCCAAAGATTTAATGGGAGCGGTGCACAGCGGTGCAGTCTGTTGGCGAAGT 1800

eGFPc.e.unc53sma

C.e.unc53 sma

K D Q K K L E Q L P T S I M P P A V S K L P S P R V A T S A T A S

GCAACTAACCCAAATTCCAACCTTCCACAAATGTCAACATCCAGGCTTCAGACTCCACAGTCAAGAATATCGAAAATTGATTCATCAAAGATTGGTATCA
CGTTGATTGGGTTTAAGGTTGAAAGGTGTTTACAGTTGTAGGTCGGAAGTCGAGGTGTCAGTTCTTATAGCTTTTAACTAAGTAGTTTCTAACCATAGT 1900

eGFPc.e.unc53sma

C.e.unc53 sma

A T N P N S N F P Q M S T S R L Q T P O S R I S K I D S S K I G I

AGCCAAAGACGCTGGACTTAAACCACCTCATCATCAACCACTTCATCAATAATACAAATTCATTCGTCGTCGAGCCGTTGAGTGGAATAATAA
TCGGTTTCTGCAGACCTGAATTTGGTGGGAGTAGTAGTTGGTGAAGTAGTTATTATGTTTAAGTAAGGCAGGCAGCTCGGCAAGCTCACCGTTATTATT 2000

eGFPc.e.unc53sma

C.e.unc53 sma

K P K T S G L K P P S S S T T S S N N T N S F R P S S R S S G N N N

TGTTGGCTCGACGATATCCACATCTGCGAAGAGCTTAGAATCATCATCAACGTACAGCTCTATTTGGAATCTAAACCGACCTACC TCCCAACTCCAAAA
ACAACCGAGCTGCTATAGGTGATAGCGCTTCTCGAATCTTAGTAGTAGTTGCATGTCGAGATAAAGCTTAGATTGGCTGGATGGAGGGTTGAGGTTTTT 2100

eGFPc.e.unc53sma

C.e.unc53 sma

V G S T I S T S A K S L E S S S T Y S S I S N L N R P T S Q L Q I

CCTTCTAGACCACAAACCCAGCTAGTTCTGTTGCTACAAC TACAAAAATCGGAAGCTCAAAGCTAGCCGCTCCGAAAGCCGTGAGCACCCCAAACTT
GGAAGATCTGGTGTGTTGGGTCGATCAAGCACACGATGTTGATGTTTTAGCCTTCGAGTTTCGATCGGCGAGGCTTTCGGCAC TCGTGGGTTTTTSAAC 2200

eGFPc.e.unc53sma

C.e.unc53 sma

P S R P Q T Q L V R V A T T T K I G S S K L A A P K A V S T P K L

CTTCTGTGAAGACTATTGGAGCAAAACAAGAGCCCGATAACAGCGGTGGTGGTGGTGGTGAATGCTGAAATTAAGTTATTTCAGTAGCAAAAACCCATC
GAAGACACTTCTGATAACCTCGTTTTGTTCTCGGGCTATTGTCGCCACCACCACCACCTTACGACTTTAATTTCAATAAGTCATCGTTTTTGGGTAG 2300

eGFPc.e.unc53sma

C.e.unc53 sma

A S V K T I G A K O E P D N S G G G G G G M L K L K L F S S K N P S

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fig 31 pEGFPsma (1 > 6960) Site and Sequence

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TTCTCATCGAATAGCCACAACCTACGAGAAAGCGGCGGCGTGCC TCAACAACAACTTTGTGCAAAATCGCTGCCCCAGTGAAAAGTGGCCTGAAG
AAGGAGTAGCTTATCGGGTGTGGATGCTCTTTCCGCCGCCGCCACGGAGTTGTTGTTGAAACAGC TTTAGCGACGGGGTCACTTTACCAGGACTTC
eGFPc.e.unc53sma
S S S N S P Q P T R K A A A V P Q Q Q T L S K I A A P V K S G L I
CCGCCGACCAGTAAGCTGGGAAGTGCCACGCTATGTGGAAGCTTTGTACGCCAAAGTTTCCTACCGTAAACGGACGCCCAATCATATCTCAACAAG
GGCGGCTGGTCAATTCGACCCCTTACGGTGCAGATACAGCTTCGAAACATGCGGTTTCAAAGGATGGCATTGCTGCGGGGTTAGTATAGAGTTGTT
eGFPc.e.unc53sma
P P T S K L G S A T S M S K L C T P K V S Y R K T D A P I I S Q Q
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D S K R C S K S S E E E S G Y A G F N S T S P T S S S T E G S L S M
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F S T S S K S S T S D E K S P S S D D L T L N A S I V T A I R Q P
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I A A T P V S P N I I N K P V E E K P T L A V K G V K S T A K K D
CAGCTCCAGCTGTTCCGCCACGTGACACCCAGCCAACAATCGGAGTTGTTAGTCCAATTATGGCACATAAGAAGTTGACAAATGACCCCGTGATATCTGA
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P P P A V P P R D T Q P T I G V V S P I M A H K K L T N D P V I S E
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fig 31 pEGFPsma (1 > 8980) Site and Sequence

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300

eGFPc.e.unc53sma

C.e.unc53 sma

K P E P E K L Q S M S I D T T D V P P L P P L K S V V P L K M T S

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310

eGFPc.e.unc53sma

C.e.unc53 sma

I R Q P P T Y D V L L K Q G K I T S P V K S F G Y E Q S S A S E D

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320

eGFPc.e.unc53sma

C.e.unc53 sma

S I V A H A S A Q V T P P T K T S G N H S L E R R M G K N K T S E S

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330

eGFPc.e.unc53sma

C.e.unc53 sma

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340

eGFPc.e.unc53sma

C.e.unc53 sma

P D N F E D S S S L S S G I S D N N E L D D I S T D D L S G V D P

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350

eGFPc.e.unc53sma

C.e.unc53 sma

A T V A S K H S D Y S H F V R H P T S S S S K P R V P S R S S T S V

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fig 31 pEGFPsma (1 > 6960) Site and Sequence

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360

-----aGFPC.e.unc53sma-----

-----C.e.unc53 sma-----

D S R S R A E Q E N V Y K L L S Q C R T S Q R G A A A T S T F G Q

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370

-----aGFPC.e.unc53sma----->

-----C.e.unc53 sma----->

H S L R S P G S T G S R . L I I I S H T T F V E V L L A L K N L P

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380

H L P L N L K H K M N A I V V V N L F I A A Y N G Y K . S N S I T N

TTTCACAAATAAGCATTTTTTCTACGCTTCTAGTGTGGTTTGTCAAACTCATCAATGTATCTTAACGCGTAAATGTAAAGCGTTAATATTTGTT
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390

F T N K A F F S L H S S C G L S K L I N V S . R V N C K R . Y F V

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400

K I R V K F L L N Q L I F . P I G R N R Q N P L . I K R I D R D R

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410

V E C C S S L E O E S T I K E R G L Q R Q R A K N R L S G R W P T T

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CACTTGGTAGTGGGATTAGTTCAAAAAACCCAGCTCCACGGCATTTCGTGATTAGCCTTGGGATTTCCTCGGGGGCTAAATCTCGAATGCCCCCTT

420

T I T L I K F F G V E V P . S T K S E P . R E P P I . S L T G I .

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CGGGCGCTTGACCGCTCTTTCTTCCCTTCTTTCGCTTTCCTCGCCCGGATCCCGGACCGTTACATCGCCAGTGGCAGCGCATTGGTGGTGTGGG

430

A G E R G E K G R E E S E R S G R . G A G K C S G H A A R N H H T

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440

R R A . C A A T G R V R W H F S G K C A R N P Y L F I F L N T F K Y

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450

V S A H E T I T L I N A S I I L K K E E S . G G K N Q L V N V C Q

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fig 31 pEGFPsma (1 > 6960) Site and Sequence

Page 7

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AATCCCACACCTTTCAGGGGTCCGAGGGGTGCTCCGCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTCCACACCTTTCAGGGGTCCGAGGG
L G C G K S P G S P A G R S M Q S M H L N . S A T R C G K S P G S 280

CAGCAGGCAGAGTATGCAAAGCATGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACCCGCCATCCCGCCCTAACCCGCCAGTTCGGC
GTCGTCCGCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTATCAGGGCGGGATTGAGGCGGGTAGGGCGGGGATTGAGGCGGGTCAAGGC
P A G R S M Q S M H L N . S A T I V P P L T P P I P P L T P P S S A 270

CCATTCTCCGCCCATGGCTGACTAATTTTTTTTATTTATGCAAGAGGCCAGGCGCCCTCGGCTCTGAGCTATTCCAGAAGTAGTGAGGAGGCTTTTTT
GGTAAGAGGGCGGGTACCGACTGATTAATAAATAAATACGCTCCGGCTCCGGCGGAGCCGAGACTCGATAAGGTCTTCATCACTCTCCGAAAAA
H S P P H G . L I F F I Y A E A E A S A S E L F Q K . . G G F F 260

GGAGGCTAGGCTTTTGCAAAGATCGATCAAGAGACAGGATGAGGATCGTTTCGATGATTGAACAAGATGGATTGCACGCAGGTCTCCGGCGGCTTGG
CCTCCGGATCCGAAACGTTTCTAGCTAGTTCCTGTCTTCTGCTTACTCTAGCAAGCGTACTAATTTGTTCTACCTAACGTGCGTCCAAGAGGCCGGCGAACC
G G L G F C K D R S R D R M R I V S H D . T R W I A R R F S G R L 490

GTGGAGAGGCTATTCCGCTATGACTGGGCACAACAGACAATCGGCTGCTCTGATGCCCGCTGTTCCGGCTGTGACGCGAGGGGCGCCGGTCTTTTTTG
CACCTCTCCGATAAGCCGATACTGACCCGTGTGTCTGTGTAGCCGACGAGACTACGGCGGCAACAAGCCGACAGTCGCGTCCCGGGGCAAGAAAAAC
G G E A I R L . L G T T D N R L L . C R R V P A V S A G A P G S F C 500

TCAAGACCGACCTGTCCGGTGCCTGAATGAAGTCAAGACGAGGCGAGCGGCTATCGTGGCTGGCCACGACGGCGTTCTTGCGCAGCTGTGCTCGA
AGTTCGGCTGGACAGGCCACGGGACTTACTTGACGTTCTGCTCCGTCGCGCCGATAGCACCAGCGGTGCTGCCCGCAAGGAACGCGTCGACACGAGCT
O D R P V R C P E . T A R R G S A A I V A G H D G R S L R S C A R 510

CGTTGTCACTGAAGCGGGAAGGGACTGGCTGCTATTGGGCGAAGTGCCGGGCGAGGATCTCTGTGTCATCTACCTTGCTCTCCGAGAAAGTATCCATC
GCAACAGTGACTTCGCCCTTCCGTGACCGACGATAACCCGCTTACGCGCCCGTCTAGAGGACAGTAGAGTGGAAACGAGGACGGCTCTTCATAGGTAG
R C H . S G K G L A A I G R S A G A G S P V I S P C S C R E S I H 520

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TACCGACTACGTTACGCGCCGACGTATGCGAATAGGCGGATGGAGGGTAAGCTGGTGGTTCGCTTTGTAGCGTAGCTCGCTCGTGATGAGCCTACC
H G . C N A A A A Y A . S G Y L P I R P P S E T S H R A S T Y S D G 530

AAGCCGGTCTTGTCGATCAGGATGATCTGGACGAAGAGCATCAGGGGCTCGCGCCAGCCGAACGTTCGCCAGGCTCAAGGCGAGCATGCCCGACGCGCA
TTCGGCCAGAACAGCTAGTCTTACTAGACCTGCTTCTGCTAGTCCCGAGCGGGTGGCTTGACAAGCGGTCGAGTTCCGCTCGTACGGGCTGCGGT
S R S C R S G . S G R R A S G A R A S R T V R Q A Q G E H A R R R 540

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CCTAGAGCAGCACTGGGTACCCTACTGACGACGAACGGCTTATAGTACCACCTTTTACCGGCGAAAGACCTAAGTAGCTGACACCGGCCGACCCACACCGC
G S R R D P V R C L L A E Y H G G K V P L F V I H R L V P A G C G 550

GACCGCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCGCTTCTCGTGCTTTACGGTATCGCCGCTC
CTGGCGATAGTCTGTATCGCAACCGATGGGCACATAACGACTTCTGCAACCGCGCTTACCCGACTGGCGAAGGAGCACGAATGCCATAGCGGCGAG
G P L S G H S V G Y P . Y C . R A V R R M G . P L P R A L R Y R R S 560

CCGATTTCGAGCGCATCGCTTCTATCGCTTCTTGACGAGTCTTCTGAGCGGGACTCTGGGGTTCGAAATGACCGACCAAGCGACGCCAACCTGGCA
GGCTAAGCGTCGCTAGCGGAAGATAGCGGAAGAACTGCTCAAGAAGACTCGCCCTGAGACCCCAAGCTTACTGGCTGGTTCGCTGCGGTTGGACGCT
R F A A H R L L S P S . R V L L S G T L G F E M T D Q A T P N L P 570

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fig 31 pEGFPsma (1 > 6960) Site and Sequence

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TCACGAGATTTCGATTCCACCGCCGCTTCTATGAAAGGTTGGGCTTCGGAATCGTTTTCCGGGACGCCGGCTGGATGATCCTCCAGCGCGGGGATCTCA
AGTGCTCTAAAGCTAAGGTGGCGGCGGAAGATACTTTCCAACCCGAAGCCTTAGCAAAAGGCCCTGCGGCCGACCTACTAGGAGGTGCGGCCCTAGAGT 580
S R D F D S T A A F Y E R L G F G I V F R D A G W M I L Q R G D L
TGCTGGAGTTCTTCGCCACCTAGGGGAGGCTAACTGAAACACGGAAGGAGACAATACCGGAAGGAACCGCGCTATGACGGCAATAAAAGACAGAA
ACGACCTCAAGAAGCGGGTGGGATCCCCCTCGGATTGACTTTTGCCCTTCTCTGTTATGGCCTTCTTGGGCGCGATACTGCCGTTATTTTCTGTCTT 590
M L E F F A H P R G R L T E T R K E T I P E G T R A M T A I K R Q H
TAAACGCACGGTGTGGGTCGTTTGTTCATAAACCGGGGTTCCGGTCCCAGGGCTGGCACTCTGTGATACCCACCGAGACCCATTGGGGCCAATAC
ATTTTGGTGGCCACAACCCAGCAACAAGTATTTGCGCCCCAAGCCAGGGTCCCAGCGTGAGACAGCTATGGGGTGGCTCTGGGGTAACCCCGGTTATG 600
K T H G V G S F V H K R G V R S Q G V H S V D T P P R P H V G Q Y
GCCGCGTTTCTTCTTTTCCCAACCCACCCCAAGTTCGGGTGAAGGCCAGGGCTCGCAGCCAACGTGGGGCGGCAGGCCCTGCCATAGCCTCAG
CGGGCGCAAGAAGGAAAAGGGTGGGGTGGGGGTTCAAGCCACTTCGGGTCCCAGCGTCGGTTGCAGCCCCGCGTCCGGGACGGTATCGGAGTC 610
A R V S S F S P P H P P S S G E G P G L A A N V G A A G P A I A S
GTTACTCATATATACTTTAGATTGATTTAAACTTCATTTTAAATTTAAAGGATCTAGGTGAAGATCCTTTTGTATAATCTCATGACCAAAATCCCTTA
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G Y S Y I L . I D L K L H F . F K R I . V K I L F D N L M T K I P .
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TGCACCTCAAAAGCAAGGTGACTCGCAGTCTGGGCATCTTTTCTAGTTTCTAGAGAAGCTCTAGGAAAAAGACGCGCATTAGACGACGAACGTTGT 630
R E F S F H . A S D P V E K I K G S S . D P F F L R V I C C L Q T
AAAAAACCCGCTACCAGCGGTGGTTTTGTTGCCGGATCAAGAGCTACCAACTCTTTTCCGAAGGTAAC TGGCTTCAGCAGAGCGCAGATACCAATA
TTTTTGGTGGCGATGGTCGCCACCAACAAACGGCCTAGTTCTCGATGGTTGAGAAAAGGCTTCCATTGACCGAAGTCGTCGCGCTATGGTTAT 640
K K P P L P A V V C L P D Q E L P T L F P K V T G F S R A Q I P H
CTGCTCTCTAGTGTAGCCGTAGTTAGGCCACCACCTCAAGAACTCTGTAGCACCGCTACATACCTCGCTCTGCTAATCCTGTTACCAGTGGCTGCTGC
GACAGGAAGATCACATCGGCATCAATCCGGTGGTGAAGTCTTGAGACATCTGGCGGATGTATGGAGCGAGACGATTAGGACAATGGTCACCGACGAGC 650
T V L L V . P . L G H H F K N S V A P P T Y L A L L I L L P V A A A
CAGTGGCGATAAGTCGTGCTTACCGGGTTGGACTCAAGACGATAGTTACCGGATAAGCGCGAGCGTGGGCTGAACGGGGGGTTCGTGCACACAGCC
GTCACCGCTATTGACGACAGAATGGCCCAACCTGAGTTCGCTATCAATGGCCTATTCCGCGTCGCCAGCCCGACTTGCCCCCAAGCACGTTGTGGG 660
S G D K S C L T G L D S R R . L P D K A Q R S G . T G G S C T Q P
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S L E R T T Y T E L R Y L Q R E L . E S A T L P E G R K A D R Y P
TAAGCGGCAGGGTCGGAACAGGAGAGCGCACGAGGGAGCTTCCAGGGGGAACGCCTGGTATCTTTATAGTCTGTGCGGTTTCGCCACCTCTGACTTGA 680
ATTGCGCGTCCCAGCCTTGTCTCTCGGTGCTCCCTCGAAGGTCCCCCTTTCGGGACCATAGAAATATCAGGACAGCCCAAGCGGTGGAGACTGAAC
V S G R V G T G E R T R E L P G G H A V Y L Y S P V G F R H L . L E
GCGTCGATTTTGTGATGCTCGTCAGGGGGCGGAGCCTATGAAAAACGCCAGCAACCGGCCCTTTTACGGTTCCTGGCTTTTGTGGCTTTTGTGCT 690
CGCAGCTAAAAACACTACGAGCAGTCCCCCGCCTCGGATACCTTTTGGGTGCTTGGCGGAAAAATGCCAAGGACCGGAAAAACCGGAAAAACGA
R R F L . C S S G G R S L V K N A S N A A F L R F L A F C V P F A

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fig 31 pEGFPsma (1 > 6960) Site and Sequence

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• CACATGTTCTTTCC TGGTTATCCCCTGATTCTGTGGATAACCGTATTACCGCCATGCAT
GTGTACAAGAAAGGACGCAATAGGGGACTAAGACACCTATTGGCATAATGGCGGTACGTA 6960
H M F F P A L S P D S V D N R I T A M H

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fig 32 pEGFPec1 (1 > 6700) Site and Sequence

Enzymes : 72 of 146 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

Page 1

3p.

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ATCAATAATTATCATTAGTTAATGCCCCAGTAATCAAGTATCGGGTATATACCTCAAGGCGCAATGTATTGAATGCCATTACCGGGCGGACCGACTGGC
L L I V I N Y G V I S S . P I Y G V P R Y I T Y G K V P A V L T
CCCAACGACCCCGCCCATTCAGCTCAATAATGACGTATGTTCCCATAGTAACGCCAATAGGGACTTTCATTGACGTCAATGGGTGGAGTATTACGGT
GGGTGCTGGGGGGCGGTAACGTCAGTTATTACTGCATACAAGGGTATCATTCGCGTTATCCCTGAAAGGTAACGTCAGTTACCCACCTCATAAATGCCA
A Q R P P P I D V N N D V C S H S N A N R D F P L T S M G G V F T V
AAACTGCCCACTTGGCAGTACATCAAGTGTATCATATGCCAAGTACGCCCCCTATTGACGTCAATGACGGTAAATGGCCCGCCTGGCATTATGCCAGTA
TTTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTCATGCGGGGATAACGTCAGTTACTGCCATTACCGGGCGGACCGTAATACGGGTCA
N C P L G S T S S V S Y A K Y A P Y . R Q . R . M A R L A L C P V
CATGACCTTATGGGACTTTCCTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGATGCGGTTTGGCAGTACATCAATGGGCGTGG
GTACTGGAATACCTGAAAGGATGAACCGTCATGTAGATGCATAATCAGTAGCGATAATGGTACCCTACGCCAAAACCGTCATGTAGTTACCCGCACT
H D L M G L S Y L A V H L R I S H R Y Y H G D A V L A V H O V A W
TAGCGGTTTACTACGGGATTTCCTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGATGCGGTTTGGCAGTACATCAATGGGCGTGG
ATCGCCAAATGAGTGCCCTAAAGGTTAGAGGTGGGGTAACGTCAGTTACCTCAAAACAAACCGTGTTTGTAGTTGCCCTGAAAGGTTTACAGCAT
I A V . L T G I S K S P P H . R Q V E F V L A P K S T G L S K M S
ACAATCCGCCCCATTGACGCAATGGGCGGTAGGCGTGTACGGTGGGAGGTCTATATAAGCAGAGCTGGTTAGTGAACCGTCAGATCCGCTAGCGCTA
TGTGAGGCGGGGTAACGCGTTTACCCGCCATCCGCACATGCCACCTCCAGATATATTGCTCGACCAATCACTTGGCAGTCTAGGCGATCGCGAT
Q L R P I D A N G R . A C T V G G L Y K Q S V F S E P S D P L A L
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P V A T M V S K G E E L F T G V V P I L V E L D G D V N G H K F S
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eGFPC.e.unc53ec1
V S G E G E G D A T Y G K L T L K F I C T T G K L P V P W P T L V T
CACCTGACCTACGGCGTGCAGTGTTCAGCCGCTACCCGACACATGAAGCAGCAGCTTCTTCAAGTCCGCCATGCCGAAGGCTACGTCCAGGAG
GTGGGAC TGGATGCCGACGTACGAAGTCGGCGATGGGGCTGGTGTACTTCGTCTGTCTGAAGAAGTTCAGGCGGTACGGGCTTCCGATGCAGGTCTCT
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T L T Y G V Q C F S R Y P D H M K Q H D F F K S A P P E G Y V Q E
CGCACCATCTTCTTCAAGGACGACGGCAACTACAAGACCCGCGCCGAGGTGAAGTTCGAGGGCGACACCTGGTGAACCGCATCGAGCTGAAGGGCATCG
GGTGGTGAAGAAGTTCCTGCTGCCGTGTATGTTCTGGGCGCGGCTCCACTTCAAGCTCCCGCTGTGGGACCACCTGGCGTAGCTCGACTTCCCGTAGC
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P T I F F K D D G N Y K T R A E V K F E G D T L V H R I E L K G I

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Page 2

ACTTCAAGGAGGACGGCAACATCCTGGGGCACAAGCTGGAGTACAACACAACAGCCACAACGTCTATATCATGGCCGACAAGCAGAAGAACGGCATC&4
TGAAGTTCCCTCCGCGTTGAGGACCCCGTGTTCGACCTCATGTTGATGTTGTCGGTGTTCAGATATAGTACC&GCTGTTCTGCTCTTCTTCCCGTAGT- 110

eGFPC.e.unc53ec1

D F K E D G N I L G H K L E Y N Y N S H N V Y I M A D K Q K N G I I

GGTGAAC TTCAAGATCCGCCACAACATCGAGGACGGCAGCGTGCAGCTCGCCGACCCTACCAGCAGAACACCCCATCGGC&GACGGCCCGTGTCTGCTG
CCACTTGAAGTTCTAGGCGGTGTTGAGCTCCTGCGCTCGCAGCTCGAGCGGCTGGTGATGGTCTGCTTGTGGGGGTAGCCGCTGCCGGGGCAGCAGCAG 120

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V N F K I R H N I E D G S V Q L A D H Y Q Q N T P I G D G P V L L

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P D N H Y L S T O S A L S K D P N E K R D H M V L L E F V T A A G

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C.e.unc53 ed

I T L G M D E L Y K S G L R S T S N V E L I P I Y T D W A N R H L S

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C.e.unc53 ed

K G S L S K S I R D I S N D F R D Y R L V S Q L I N V I V P I N E

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F S P A F T K R L A K I T S N L D G L E T C L D Y L K N L G L D C

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C.e.unc53 ed

S K L T K T D I D S G N L G A V L Q L L F L L S T Y K Q K L R Q L I

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fig 32 pEGFPec1 (1 > 6700) Site and Sequence

Page 3

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C.e.unc53 ed
K P K T S G L K P P S S S T T S S N N T N S F R P S S R S S G N N H
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V G S T I S T S A K S L E S S S T Y S S I S N L N R P T S Q L Q I
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P S R P Q T Q L V R V A T T T K I G S S K L A A P K A V S T P K L
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A S V K T I G A K O E P D N S G G G G G G M L K L K L F S S K N P S

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S S S N S P Q P T R K A A A V P Q Q Q T L S K I A A P V K S G L A

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C.e.unc53 ed

P P T S K L G S A T S M S K L C T P K V S Y R K T D A P I I S Q Q

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C.e.unc53 ed

D S K R C S K S S E E E S G Y A G F N S T S P T S S S T E G S L S M

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H S T S S K S S T S D E K S P S S D D L T L N A S I V T A I R Q P

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C.e.unc53 ed

I A A T P V S P N I I N K P V E E K P T L A V K G V K S T A K K D

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P P P A V P P R D T Q P T I G V V S P I M A H K K L T N D P V I S E

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fig 32 pEGFPec1 (1 > 6700) Site and Sequence

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C.e.unc53 ed
K P E P E K L Q S M S I D T T D V P P L P P L K S V V P L K M T S
ATCCGACAACCACCAACGTACGATGTTCTTCTAAAACAAGGAAAAATCACATCGCCTGTCAGTCTTTGGATATGAGCAGTCGTCCGCTCTGAAGACT
TAGGC TGTGGTGGTTGCATGCTACAAGAAGATTTTGTTCCTTTTAGTGTAGCGGACAGTTCAGCAAACCTATACTCGTCAGCAGGCGCAGACTTCTGA
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C.e.unc53 ed
I R O P P T Y D V L L K Q G K I T S P V K S F G Y E Q S S A S E D
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C.e.unc53 ed
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H I C R G F T C F K K P P T P P P E P E T . N E C N C C C . L V Y C
CAGCTTATAATGGTTACAAATAAAGCAATAGCATCACAAATTTACAAATAAAGCATTTTTTTCAC TGCATTC TAGTTGTGGTTTGCCAAACTCATCA
GTGGAATATTACCAATGTTTATTTTCGTTATCGTAGTGTITAAAGTGTITATTTTCGTAAGGAGTACGTAAGATCAACACCAACAGGTTTGAGTAGTT
S L . V L Q I K O . H H K F H K . S I F F T A F . L V F V O T H Q

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C I L T R K L . A L I F C . N S R . I F V K S A H F L T N R P K S
CAAAATCCCTTATAAATCAAAGAATAGACCGAGATAGGGTTGAGTGTGTTCAGTTTGAACAAGAGTCCACTATTAAGAACGTGGACTCCAACGTC
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A K S L I N O K N R P R . G . V L F Q F G T R V H Y . R T W T P T S
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K G E K P S I R A M A H Y V N H P N O V F W G R G A V K H . I G
ACCTAAAGGGAGCCCCGATTTAGAGCTTGACGGGGAAAGCCGGCGAAGTGGCGAGAAAGGAAGGAAGCAAGGAGCGGGCGCTAGGGCGCT
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T L K G A P D L E L D G E S R R T V R E R K G R K R K E R A L G R
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CCGTTACATCGCCAGTGGCAGCGCATTTGGTGGTGTGGGCGGCGCAATTACGCGCGATGTCCCGCGCAGTCCACCGTGAAGAGCCCTTTACACGGC 410
V O V . R S R C A . P P H P P R L M R R Y R A R O V A L F G E M C A
GGAACCCCTATTTGTTTATTTTCTAAATACATTCAAATATGTATCCGCTCATGAGACAATAACCTGATAAATGCTTCAATAATATTGAAAAAGGAAGA
CCTTGGGGATAAAACAAATAAAAGATTATGTAAGTTTATACATAGGCGAGTACTCTGTTATTGGGACTATTTACGAAGTTATTATACTTTTCTTCT 420
E P L F V Y F S K Y I O I C I R S . D N N P D K C F N N I E K G R
GTCCTGAGGCGGAAGAACCAGCTGTGGAATGTGTGTCAGTTAGGGTGTGGAAGTCCCAGGCTCCCAGCAGGCAGAGATGCAAGCATGCATCTC
CAGGACTCCGCTTTCTTGGTGCACACCTTACACACAGTCAATCCACACCTTTAGGGGTCCGAGGGGTGCTCCGCTTTCATACGTTTCTGACGTAGAG 430
V L R R K E P A V E C V S V R V V K V P R L P S R O K Y A K H A S
AATTAGTCAGCAACCAGGTGTGGAAGTCCCAGGCTCCCAGCAGGCAGAGTATGCAAGCATGCATCTCAATTAGTCAGCAACCATAGTCCCGCCC
TTAATCAGTCGTTGGTCCACACCTTTAGGGGTCCGAGGGGTGCTCCGCTTTCATACGTTTCGTACGTAGAGTAAATCAGTCGTTGGTATCAGGGCGGG 440
Q L V S N Q V W K V P R L P S R O K Y A K H A S Q L V S N H S P A P
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ATTGAGGCGGGTAGGGCGGGATTGAGGCGGGTCAAGGCGGGTAAGAGGCGGGTACCGACTGATTAATAAAAAATAAATACGCTCCGGCTCCGGCGGAG 450
N S A H P A P N S A Q F R P F S A P V L T N F F Y L C R G R G R L
GGCCTCTGAGCTATTCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCTAGGCTTTTGAAGATCGATCAAGAGACAGGATGAGGATCGTTTCGCATGAT
CCGAGACTCGATAAGGTCTTCATCACTCTCCGAAAAAACCTCCGATCCGAAAAAGTTCAGTCTAGTCTCTGCTACTCCAGCAAGCGTACTA 460
G L . A I P E V V R R L F W R P R L L Q R S I K R Q D E D R F A .
TGAACAAGATGGATTGCACGCAGGTTCTCCGGCCGCTTGGGTGGAGAGGCTATTCCGCTATGACTGGGCAACAACAGACAATCGGCTGCTCTGATGCCGC
ACTTGTCTTACCTAACGTGCGTCCAAGAGGCGGCGAACCACCTCTCCGATAAGCGGATGACCCGTTGTTCTGTTAGCCGACGAGACTACGGCGG 470
L N K M D C T O V L R P L G V R G Y S A M T G H N R O S A A L M P P
GTGTCCGGCTGTACAGCGAGGGGCGCCGGTCTTTTTGTCAAGACCGACCTGTCCGGTGGCCGTAATGAACGCAAGACGAGGACGCGGCTATCGT
CACAAAGGCGACAGTCCGCTCCCGCGGGCAAGAAAAACAGTTCTGGCTGGACAGGCCAGGGACTTACTTGACGTTCTGCTCCGTCGCGCGGATAGGA 480
C S G C O R R G A R F F L S R P T C P V P . M N C K T R O R G Y F

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GGC TGGCCACGACGGGCGTTCTTGCACAGCTGTGCTCGACGTTGCTACTGAAGCGGGAAGGGACTGGCTGCTATTGGGCGAAGTGCCGGGGCAGGATCT
CCGACCGGTGCTGCCCGCAAGGAACGCGTCGACACGAGCTGCAACAGTGACTTCGCCCTTCCTGACCGACGATAACCCGCTTCACGGCCCCGCTCTAGA
G V P R R A F L A Q L C S T L S L K R E G T G C Y V A K C R G R I
CCTGTCTCTCACCTTGCTCTGCGGAGAAAGTATCCATCATGGCTGATGCAATGCGGCGGCTGCATACGCTTGATCCGGCTACCTGCCCATTCGACCAC
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S C H L T L L L P R K Y P S V L M Q C G G C I R L I R L P A H S T I
CAAGCGAAACATCGCATCGAGCGAGCAGTACTCGGATGGAAGCGGTCTTGTGCTATCAGGATGATCTGGACGAAGAGCATCAGGGGCTCGCGCCAGCCG
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K R N I A S S E H V L G W K P V L S I R M I V T K S I R G S R Q P
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G L T A S S C F T V S P L P I R S A S P S I A F L T S S S E R D S
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G V R N D R P S D A Q P A I T R F R F H R R L L . K V G L R N R F
CGGGACCGCGCTGGATGATCTCCAGCGCGGGGATCTCATGCTGGAGTTCTTCGCCCACCTAGGGGGAGGCTAACTGAAACACGGAAGGAGACAATAC
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P G R R L D D P P A R G S H A G V L R P P . G E A N . N T E G D N I
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G R N P R Y D G N K K T E . N A R C V V V C S . T R G S V P G L A
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L C R Y P T E T P L G P I R P R F F L F P T P P P K F G . R P R A
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R S Q R R G G R P C H S L R L L I Y T L D . F K T S F L I . N D L G
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E O P F . . S H D Q N P L T . V F V P L S V R P R R K D Q R I F L

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R S F F S A R N L L L A N K K T T A T S G G L F A G S R A T N S F
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CGTCGCCAGCCGACTTGCCCCCAAGCAGTGTGTCGGGTCGAACCTCGCTTGTGATGTGGCTTGACTCTATGGATGTCGCACTCGATACTCTTTCCG 3400
A A V G L N G G F V H T A Q L G A N D L H R T E I P T A . A M R K
GCCACGCTTCCC GAAGGGAGAAAGGCGGACAGGTATCCGGTAAGCGGCAGGGTCGGAACAGGAGAGCGCACGAGGGAGCTTCCAGGGGGAAACGCC TGGT
CGGTGCGAAGGGCTTCCCTCTTTCCGCC TGTCCATAGGCCATTCCGCGTCCAGCCTTGTCTCTCGCGTGCTCCCTCGAAGGTCCCCCTTTGCGGACCA 3500
R H A S R R E K G G Q V S G K R O G R N R R A H E G A S R G K R L V
ATCTTTATAGTCTGTGCGGTTTCGCCACCTCTGACTTGAGCGTCGATTTTGTGATGCTCGTCAGGGGGGCGGAGCCTATGGAAAAACGCCAGCAACGC
TAGAAATATCAGGACAGCCCAAAGCGGTGGAGACTGAAC TCGCAGCTAAAAACACTACGAGCAGTCCCCCGCCTCGGATACCTTTTTCGCGTCTGTGCG 3600
S L . S C R V S P P L T . A S I F V M L V R G A E P M E K R Q Q R
GGCCTTTTACGGTTCCTGGCCTTTTGTGCTCACATGTTCTTTCTGCGTTATCCCTGATTCTGTGGATAACCGTATTACCGCCATGCAT
CCGGAAAAATGCCAAGGACCGGAAACGACCGGAAACGAGTGTACAAGAAAGGACGCAATAGGGGACTAAGACACCTATTGGCATAATGGCGGTACGT 3700
G L F T V P G L L L A F C S H V L S C V I P . F C G . P Y Y R H A

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fig 33 pEGFPxba (1 > 5447) Site and Sequence

Enzymes : 72 of 146 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

Page 1

TAGTTATTAATAGTAATCAATTACGGGGTCATTAGTTTCATAGCCCATATATGGAGTTCGCGTTACATAACTTACGGTAAATGGCCCGCTGGCTGACCG
ATCAATAATTATCATTAGTTAATGCCCCAGTAATCAAGTATCGGGTATATACCTCAAGGCGCAATGTATTGAATGCCATTTACGGGGCGGACCGACTGGC
L L I V I N Y G V I S S . P I Y G V P R Y I T Y G K V P A V L T
CCCAACGACCCCGCCCATTTGACGTCAATAATGACGTATGTTCCCATAGTAACGCCAATAGGGACTTTCCATTGACGTCAATGGGTGGAGTATTTACGGT
GGGTTGCTGGGGGCGGGTAACGCGATTATTACTGCATACAAGGGTATCATTGCGGTTATCCCTGAAAGGTAACGCGATTACCCACCTCATAAATGCCA
A Q R P P P I D V N N D V C S H S N A N R D F P L T S M G G V F T V
AAACTGCCCACTTGGCAGTACATCAAGTGTATCATATGCCAAGTACGCCCCCTATTGACGTCAATGACGGTAAATGGCCCGCTGGCATTATGCCAGTA
TTTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTCATCGGGGGGATAACTGCGATTACTGCCATTACCGGGCGGACCGTAATACGGGTCA
N C P L G S T S S V S Y A K Y A P Y . R Q . R . M A R L A L C P V
CATGACCTTATGGGACTTTCCTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGATGCGGTTTTGGCAGTACATCAATGGGCGTGGA
GTACTGGAATACCCTGAAAGGATGAACCGTCATGTAGATGCATAATCAGTAGCGATAATGGTACCCTACGCCAAAACCGTCATGTAGTTACCCGACCT
H D L M G L S Y L A V H L R I S H R Y Y H G D A V L A V H O V A V
TAGCGGTTTGAATCAGGGGATTTCGAAGTCTCCACCCCATTTGACGTCAATGGGAGTTTGTGTTTGGCACCAAAATCAACGGGACTTTCCAAAATGTCGTA
ATCGCCAAATGAGTGCCCTTAAAGGTTTCAAGGTTGGGTAACGCGATTACCTCAACAAAACCGTGGTTTTAGTTGCCCTGAAAGGTTTTACAGCAT
I A V . L T G I S K S P P H . R Q V E F V L A P K S T G L S K M S
ACAACCTCGCCCCATTGACGCAATGGGCGGTAGGCGGTACGGTGGGAGGTCTATATAAGCAGAGCTGGTTTTAGTGAACCGTCAGATCCGCTAGCGCTA
TGTTGAGGCGGGTAACGCGTTTACCCGCCATCCGCACATGCCACCTCCAGATATATTGCTCTGACCAAAATCACTTGGCAGTCTAGGCGATCGCGAT
D L R P I D A N G R . A C T V G G L Y K Q S W F S E P S D P L A L
CGCGTCGCCACCATGGTGAGCAAGGGCGAGGAGCTGTTACCGGGGTGGTGGCCATCCTGGTCGAGCTGGACGGCGACGTAAACGGCCACAGTTACGCG
GGCCAGCGGTGGTACCACTCGTTCCCGCTCCTCGACAAGTGGCCCCACCACGGGTAGGACCAGCTCGACCTGCCGCTGCATTGGCCGGTGTCAAGTGGC
eGFPC.e.unc53xba
P V A T M V S K G E E L F T G V V P I L V E L D G D V N G H K F S
TGTCGGGCGAGGGCGAGGCGATGCCACCTACGGCAAGCTGACCGTGAAGTTTCATCTGCACCACCGGCAAGCTGCCCGTGCCCTGGCCCAACCTCGTGAC
ACAGGCGCGTCCCGCTCCCGCTACGGTGGATGCCGTTTCGACTGGGACTTCAAGTAGACGTGGTGGCCGTTTCGACGGGCACGGGACCGGGTGGGAGCACTC
eGFPC.e.unc53xba
V S G E G E G D A T Y G K L T L K F I C T T G K L P V P W P T L V T
CACCTGACCTACGCGGTGCGAGTGTTCAGCCGCTACCCCGACCATGAAGCAGCACGACTTCTTCAAGTCCGCCATGCCCGAAGGCTACGTCCAGGAG
GTGGGACTGGATGCCGACGTACGAAGTCGGCGATGGGGCTGGTGTACTTCGTCGTCTGAAGAAGTTCAGGCGGTACGGGCTTCCGATGCAGGTCTCT
eGFPC.e.unc53xba
T L T Y G V O C F S R Y P D H M K Q H D F F K S A M P E G Y V Q E
CGCACCATCTTCTTCAAGGACGACGGCAACTACAAGACCCGCGCGAGGTGAAGTTCGAGGGCGACACCTGGTGAACCGCATCGAGCTGAAGGGCATCG
GCCTGGTAGAAGAAGTTCCTGCTGCCGTTGATGTTCTGGGCGCGGCTCCACTTCAAGCTCCCGCTGTTGGGACCACTTGGCGTAGCTCGACTTCCCGTAGC
eGFPC.e.unc53xba
R T I F F K D D G N Y K T R A E V K F E G D T L V N R I E L K G I

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fig 33 pEGFPxba (1 > 5447) Site and Sequence

Page 1

ACTTCAAGGAGGACGGCAACATCTGGGGCACAAAGCTGGAGTACAACACAAGCCACAACGTCTATATCATGGCCGACAAGCAGAAGAACGGCATCAA
TGAAGTTCTCTGCGGTTGTAGGACCCCGTGTTCGACCTCATGTTGATGTTGTCGGTGTTCAGATATAGTACCGGCTGTTCTGCTCTTGGCCGTAGTT

110

eGFPC.e.unc53xba

D F K E D G N I L G H K L E Y N Y N S H N V Y I M A D K Q K N G I

GGTGAACCTCAAGATCCGCCACAACATCGAGGACGGCAGCGTGCAGCTCGCCGACCACTACCAGCAGAACACCCCATCGGCACGGCCCCGTGCTGCTG
CCACTTGAAGTTCTAGGCGGTGTGTAGCTCTGCGCTCGCAGCTCGAGCGGCTGGTGTGTTGCTGCTTGTGGGGGTAGCCGCTGCCGGGGCACGACGAC

120

eGFPC.e.unc53xba

V N F K I R H N I E D G S V Q L A D H Y Q Q N T P I G D G P V L L

CCCGACAACCACTACCTGAGCACCCAGTCCGCCCTGAGCAAGACCCCAACGAGAAGCGCGATCACATGGTCTCTGCTGGAGTTCGTGACCGCCGCCGGGA
GGGCTGTGGTGTGAGTCTGTTGGTTCAGGCGGGACTCGTTTCTGGGGTTGCTCTTCGCGCTAGTGATACAGGACGACCTCAAGCACTGGCGGGCGCCCT

130

eGFPC.e.unc53xba

P D N H Y L S T Q S A L S K D P N E K R D H M V L L E F V T A A G

TCACTCTCGCATGGACGAGCTGTACAAGTCCGGACTCAGATCTACGTCAAATGTAGAATTGATACCAATCTACACGGATTGGGCCAATGGCACCTTTC
AGTGAGAGCCGTACCTGCTCGACATGTTCAAGGCTGAGTCTAGATGCAGTTTACATCTTAACATATGGTTAGATGTGCTTAACCCGGT TAGCGTGGAAAG

140

eGFPC.e.unc53xba

C.e.unc53 xba

I T L G M D E L Y K S G L R S T S N V E L I P I Y T D V A N R H L S

GAAGGGCAGCTTATCAAAGTCGATTAGGGATATTTCCAATGATTTTCGCGACTATCGACTGGTTTCTCAGCTTATTAATGTGATCGTTCCGATCAACGAA
CTTCCCGTCGAATAGTTTCAGCTAATCCCTATAAAGGTTACTAAAGCGCTGATAGCTGACCAAGAGTCGAATAATTACAC TAGCAAGGCTAGTTGCTT

150

eGFPC.e.unc53xba

C.e.unc53 xba

K G S L S K S I R D I S N D F R D Y R L V S Q L I N V I V P I N E

TTCTCGCTGCATTACGAAACGTTTGGCAAAATCACATCGAACCTGGATGGCTCGAAACGTGTCTCGACTACCTGAAAAATCTGGGTCTCGACTGCT
AAGAGCGGACGTAAGTGCTTTTGCAACCGTTTTAGTGAGCTTGGACCTACCGGAGCTTTGCACAGAGCTGATGGACTTTTAGACCCAGAGCTGACGA

160

eGFPC.e.unc53xba

C.e.unc53 xba

F S P A F T K R L A K I T S N L D G L E T C L O Y L K N L G L D C

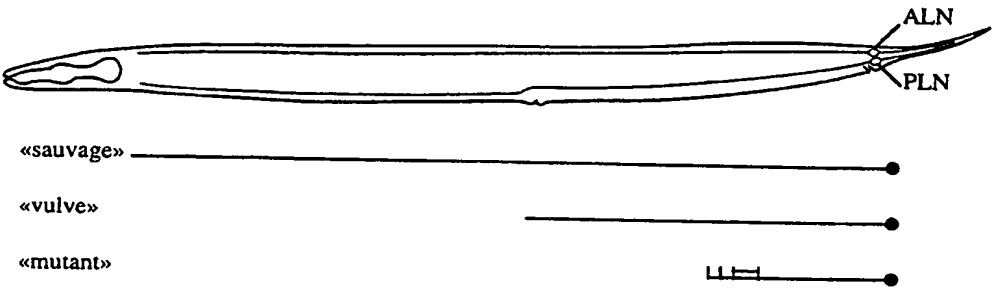
CGAAACTACCAAAACCGATATCGACAGCGGAAACTTGGGTGCAGTTCTCCAGCTGCTCTCTCTGCTCTCCACCTACAAGCAGAAGCTTCGGCAACTGAA
GCTTTGAGTGGTTTTGGCTATAGCTGTGCGCTTTGAACCCACGTCAAGAGGTCGACGAGAAGGACGAGAGGTGGATGTTCTGCTTTCGAAGCCGTGACTT

170

eGFPC.e.unc53xba

C.e.unc53 xba

S K L T K T D I O S G N L G A V L O L L F L L S T Y K Q K L R O L I



| souches | phénotypes | sauvage | vulve | mutant | nombre |
|--------------------------------------------------------------|------------|---------|-------|--------|--------|
| wt | ALN | 100 | 0 | 0 | 70 |
| | PLN | 100 | 0 | 0 | 70 |
| <i>unc-53(n152)</i> | ALN | 0 | 26 | 74 | 50 |
| | PLN | nd | nd | nd | |
| <i>unc-53(n152)</i> <i>pAΔunc-53</i> | ALN | 95 | 0 | 5 | 65 |
| | PLN | 77 | 0 | 23 | 44 |
| <i>unc-53(n152)</i> <i>pAΔunc-53-H1</i> | ALN | 25 | 60 | 15 | 56 |
| | PLN | 52 | 16 | 32 | 25 |

Figure 51b

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fig 52 pLM5 (1 > 5425) Site and Sequence

Page 1

Enzymes : All 148 enzymes (No Filter)

Settings: Linear, Certain Sites Only, Standard Genetic Code

GACGGATCGGGAGATCTCCCGATCCCCATGGTGCAGTCTCAGTACAATCTGCTCTGATGCCGCATAGTTAAGCCAGTATCTGCTCCCTGCTTGTGTGT
CTGCC TAGCCCTCTAGAGGGCTAGGGGATACCAGCTGAGAGTCATGTTAGACGAGACTACGGCGTATCAATTCGGTTCATAGACGAGGGACGAACACACA
T D R E I S R S P M V D S Q Y N L L . C R I V K P V S A P C L C V 100

GGAGGTCGCTGAGTAGTGC CGAGCAAAATTTAAGCTACAACAAGGCAAGGCTTGACCGACAATTGCATGAAGAATCTGCTTAGGGTTAGGCGTTTTGCG
CCTCCAGCGACTCATCACGCGCTCGTTTTAAATTCGATGTTGTTCCGTTCCGAAGTGGCTGTTAAGCTACTTCTTAGACGAATCCCAATCCGCAAAACGC
G G R . V V R E O N L S Y N K A R L D R O L H E E S A . G . A F C 200

CTGCTTCGCGATGACGGGCAGATATACGCGTTGACATTGATTATTGACTAGTTATTAATAGTAATCAATTACGGGGTCATTAGTTCATAGCCCATATA
GACGAAGCGCTACATGCCCGTCTATATGCGCAACTGTAACATAAATGATCAATAATTATCATTAGTTAATGCCCGAGTAATCAAGTATCGGGTATAT
A A S R C T G Q I Y A L T L I I D . L L I V I N Y G V I S S . P I Y 300

TGGAGTTCGCGTTACATAACTTACGGTAAATGGCCCGCTGGCTGACCGCCCAACGACCCCGCCCATTGACGTCAATAATGACGTATGTTCCCATAGT
ACCTCAAGGCGCAATGTATTGAATGCCATTACCGGGCGGACCGACTGGCGGGTGTCTGGGGGCGGGTAACGTCAGTTATTACTGCATACAAGGGTATCA
G V P R Y I T Y G K V P A V L T A Q R P P P I D V N N D V C S H S 400

AACGCCAATAGGGACTTTCCATTGACGTCAATGGGTGGACTATTTACGGTAAACTGCCCACTTGGCAGTACATCAAGTGTATCATATGCCAAGTACGCCC
TTGCGGTTATCCCTGAAAGGTAAGTGCAGTTACCCACCTGATAAATGCCATTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTCATGCGGG
N A N R D F P L T S M G G L F T V N C P L G S T S S V S Y A K Y A 500

CCTATTGACGTCAATGACGGTAAATGGCCCGCTGGCATTTATGCCAGTACATGACCTTATGGGACTTTCTACTTGGCAGTACATCTACGTATTAGTCA
GGATAACTGCAGTTACTGCCATTTCACGGGCGGACCGTAATACGGGTGATGTACTGGAATACCTGAAAGGATGAACCGTCATGTAGATGCATAATCAGT
P Y . R Q . R . M A R L A L C P V H D L M G L S Y L A V H L R I S H 600

TCGCTATTACCATGGTGATGCGGTTTTGGCAGTACATCAATGGGCGTGGATAGCGGTTTGACTCAGGGGATTTCAGGCTCCACCCCATGACGTCAA
AGCGATAATGGTACCACTACGCCAAAACCGTCATGTAGTTACCCGCACCTATCGCCAAACTGAGTGCCCTTAAAGGTTACAGAGTGGGGTAACGTCAGTT
R Y Y H G D A V L A V H Q W A V I A V . L T G I S K S P P H . R O 700

TGGGAGTTTGTGTTTGGCACCAAAATCAACGGGACTTTCCAAAATGTCGTAACAACTCGCCCCATTGACGCAATGGGCGGTAGGCGGTGACGGTGGGAG
ACCTCAAAACAAACCGTGGTTTTAGTTGCCCTGAAAGGTTTACAGCATGTTGAGGCGGGTAACGCGTTTACCCGCCATCCGCACATGCCACCCCT
W E F V L A P K S T G L S K M S . Q L R P I D A N G R . A C T V G 800

GTCTATATAAGCAGAGCTCTCTGGCTAACTAGAGAACCCTGCTTACTGGCTTATCGAAATTAATACGACTCACTATAGGGAGACCCAAGCTGGCTAGC
CAGATATATTCGCTTCGAGAGACCGATTGATCTCTTGGGTGACGAATGACCGAATAGCTTTAATTATGCTGAGTGATATCCCTCTGGGTTGACCGGATCG
G L Y K O S S L A N . R T H C L L A Y R N . Y D S L . G D P S V L A 900

GTTTAACTTAAGCTTACCATGGGGGTTCTCATCATCATCATCATGATGGTATGGCTAGCATGACTGGTGGACAGCAATGGGTCGGGATCTGTACGAC
CAAATTTGAATTCGAATGGTACCCCAAGAGTAGTAGTAGTAGTAGTACCATACCGATCGTACTGACCACTGTCTGTTTACCCAGCCCTAGACATGCTG
F K L K L T M G G S H H H H H H G M A S M T G G Q Q M G R D L Y D 1000

T7 promoter priming site

ProBond binding domain

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fig 52 pLM5 (1 > 5425) Site and Sequence

Page 2

GATGACGATAAGGTACCTAGGATCCATGCAAAATGAGGAGGAGGAGCCAGAGAAGAAGGAGGTATCGGAGCTGCGCTCTGAGCTATGGGAGAAGGAAATGA
CTACTGCTATTCCATGGATCCTAGGTACGTTTACTCCTCCTCCTCGGTCTCTTCTTCTCCATAGCCTCGACGCGAGACTCGATACCCTCTTCTTTACT
pLM5 insert = U3
D D D K V P R I H A N E E E E P E K K E V S E L R S E L V E K E N
AGCTTACAGACATCCGCTTGGAGGCCCTCAACTCTGCCACCAACTGGATCAGCTTCGGGAGACCATGCACAACATGCAGTTGGAGGTGGACCTGCTGA
TCGAATGCTGTAGGCGAACCTCCGGGAGTTGAGACGGGTGGTTGACCTAGTCGAAGCCCCTGGTACGTGTTGTACGTCAACCTCCACCTGGACGACT
pLM5 insert = U3
ORF U3
K L T D I R L E A L N S A H Q L D Q L R E T M H N M Q L E V D L L A
AGCAGAGAATGACCGACTGAAGGTAGCCCCAGGCCCTCATCAGGCTCCACTCCAGGGCAGGTCCCTGGATCATCTGCATTATCTTCCCACGCCGCTCC
TCGTCTCTTACTGGCTGACTTCATCGGGGTCCGGGGAGTAGTCCGAGGTGAGGTCCCGTCCAGGGACCTAGTAGACGTAATAGAAGGGGTGCGCGGAGG
pLM5 insert = U3
ORF U3
A E N D R L K V A P G P S S G S T P G Q V P G S S A L S S P R R S
CTAGGCCCTGGCACTACCCATTCCTTCGGCCCCAGTCTTGAGACACAGACCTGTACCCATGGATGGCATCAGTACTTGTGGTCCAAAGGAGGAAGTGA
GATCCGGACCGTGAGTGGGTAAGGAAGCCGGGGTCAGAACGCTCTGTGTCTGGACAGTGGGTACCTACCGTAGTCATGAACACCAGGTTTCTCTCTTCA
pLM5 insert = U3
ORF U3
L G L A L T H S F G P S L A D T D L S P M D G I S T C G P K E E V
CCCTCCGGGTGGTGGTGAAGATGCCCCSCAGCACATCATCAAGGGGACTTGAAGCAGCAGGAATTCTTCTGGGCTGTAGCAAGGTCAGTGGAAAAGT
GGGAGGCCACCACCACTCCTACGGGGCGTCTGTAGTAGTTTCCCCTGAACCTCGTCTCTTAAGAAGGACCCGACATCGTTCCAGTCACCTTTTCA
pLM5 insert = U3
ORF U3
T L R V V V R M P P Q H I K G D L K Q Q E F F L G C S K V S G V V
TGACTGGAAGATGCTGGATGAAGCTGTTTCCAAGTGTTCAGGACTATATTTCTAAAATGGACCCAGCCTCTACCTGGGACTAAGCACTGAGTCCATC
ACTGACCTTCTACGACCTACTTCGACAAAAGGTTCAAGTTCCTGATATAAAGATTTTACC TGGGTCGGAGATGGGACCTGATTCGTGACTCAGGTAG
pLM5 insert = U3
ORF U3
D V K M L D E A V F Q V F K D Y I S K M D P A S T L G L S T E S I

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fig 52 pLM5 (1 > 5425) Site and Sequence

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```
CATGGCTACAGCATCAGCCACGTGAAACGAGTGTGGATGCAGAGCCCCCGAGATGCCTCCTTGCCGTCGAGGTGTCAATAACATATCAGTCTCCCTCA
GTACCGATGTCGTAGTCGGTGCACTTTGCTCACAACCTACGTCTCGGGGGGCTCTACGGAGGAACGGCAGCTCCACAGTTATTGTATAGTCAGAGGGAG
700

pLM5 insert = U3

ORF U3
H G Y S I S H V K R V L D A E P P E M P P C R R G V N N I S V S L

AASGTCTGAAGGAGAAATGCGTCGACAGCCTGGTGTTCGAGACGCTGATCCCAAGCCGATGTCAGCACTACATAAGCCTCCTGCTGAAGCACCGGGG
TTCAGACTTCCTCTTTACGCAGCTGTCGGACCACAAGCTCTGCGACTAGGGGTTCGGCTACTACGTCGTGATGTATTTCGGAGGACGACTTCGTGGCCGC
1200

pLM5 insert = U3

ORF U3
K G L K E K C V D S L V F E T L I P K P M M Q H Y I S L L L K H R R

CCTCGTCTCTCGGGCCCCAGCGGCACGGGCAAGACCTACCTGACCAATCGCTTGGCCGAGTACCTGGTGGAGCGCTCTGGCCGTGAGGTCACAGAGGGG
GGAGCAGGAGAGCCCGGGTGGCCGTGCGCGTCTGGATGGACTGGTTAGCGAACCGGCTCATGGACCACCTCGCGAGACCGGCACCTCCAGTGTCTCCCS
1300

pLM5 insert = U3

ORF U3
L V L S G P S G T G K T Y L T N R L A E Y L V E R S G R E V T E G

ATCGTCAGCACCTTCAACATGCACCAGCAGTCTTGCAAGGATCTGCAACTGTATCTTTCCAACCTAGCCAACCAGATAGACCGGGAACAGGAATTGGGG
TAGCAGTCGTGGAAGTGTACGTGGTCGTCAGAACGTTCTAGACGTTGACATAGAAAGGTTGGATCGGTTGGTCTATCTGGCCCTTGTCTTAACCCG
2000

pLM5 insert = U3

ORF U3
I V S T F N M H Q Q S C K D L Q L Y L S N L A N O I D R E T G I G

ATGTGCCCTGGTGATTCTATTGGATGACCTGAGTGAAGCAGGCTCCATCAGTGAGTTGGTCAATGGGGCCCTCACCTGCAAGTATCATAAATGTCCTTA
TACACGGGGACCACTAAGATAACCTACTGGACTCACTTCGTCCGAGGTAGTCACTCAACCAGTTACCCCGGAGTGGACGTTTCATAGTATTACAGGGA
3100

pLM5 insert = U3

ORF U3
D V P L V I L L D D L S E A G S I S E L V N G A L T C K Y H K C P Y

TATTATAGGTACCACCAATCAGCCTGTAAAAATGACACCCAACCATGGCTTGCACTTGAGCTTCAGGATGTTGACCTTCTCCAACAACGTGGAGCCAGCC
ATAATATCCATGGTGGTTAGTCGGACATTTTACTGTGGGTGGTACCGAAGTGAACTCGAAGTCTTACAAC TGGAAAGAGGTTGTTGCACCTCGGTCCG
3200

pLM5 insert = U3

ORF U3
I I G T T N Q P V K M T P N H G L H L S F R M L T F S N N V E P A
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fig 52 pLM5 (1 > 5425) Site and Sequence

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AATGGCTTCCTGGTTCGTTACCTGAGGAGGAAGCTGGTAGAGTCAGACAGCGACATCAATGCCAACAAAGGAAGAGCTGCTTCGGGTGCTCGACTGGGTAC
TTACCGAAGGACCAAGCAATGGACTCCTCCTTCGACCATCTCAGTCGTGCGTGTAGTTACGGTTGTTCCCTTCGACGAAGCCCACGAGCTGACCCATG 330

pLM5 insert = U3

ORF U3

N G F L V R Y L R R K L V E S D S D I N A N K E E L L R V L D W V

CCAAGCTGTGGTATCATCTCCACACCTTCCTTGAGAAGCACAGCACCTCAGACTTCCTCATCGGCCCTTGCTTCTTCTGTCGTGTCCTTGGCAATTGA 240

GGTTCGACACCATAGTAGAGGTGTGAAGGAACCTTCGTGTCGTGGAGTCTGAAGGAGTAGCCGGGAACGAAGAAAGACAGCACAGGGTAACCGTAAC

pLM5 insert = U3

ORF U3

P K L V Y H L H T F L E K H S T S D F L I G P C F F L S C P I G I E

GGACTTCGGACCTGGTTCATTGACCTGTGGAACAACCTCTATCATTCCCTATCTACAGGAAGGAGCCAAAGGATGGGATAAAGGTCCATGGACAGAAAGCT 250

CCTGAAGGCC TGGACCAAGTAAC TGGACACCTTGTGAGATAGTAAGGGATAGATGTCCTTCCTCGGTTCTACCTATTTCAGGTACCTGCTTTTCGA

pLM5 insert = U3

ORF U3

D F R T V F I D L W N N S I I P Y L Q E G A K D G I K V H G Q K A

GCTTGGGAGGACCCAGTGAATGGGTCCGGGACACACTTCCCTGGCCATCAGCCCAACAAGACCAATCAAAGCTGTACCACCTGCCCCACCCACCGTGG 260

CGAACCTCCTGGGTACCTTACCCAGGCCCTGTGTGAAGGGACCGGTAGTCGGGTGTTCTGGTTAGTTTCGACATGGTGGACGGGGTGGGTGGCACC

pLM5 insert = U3

ORF U3

A V E D P V E V V R D T L P V P S A Q Q D Q S K L Y H L P P P T V

GCCCTCAGCAGATTGCCTCACCTCCCGAGGATAGGACAGTCAAAGACAGCACCCCAAGTTCTCTGGAATCAGATCCTCTGATGGCCATGCTGCTGAACCT 270

CGGGAGTGTGTAACGGAGTGGAGGGCTCTATCCTGTCAGTTTCTGTCGTGGGGTTCAAGAGACC TGAGTCTAGGAGAC TACCGGTACGACGACTTTGA

pLM5 insert = U3

ORF U3

G P H S I A S P P E D R T V K D S T P S S L D S D P L M A M L L K L

TCAAGAAGCTGCCAACACATTGAGTCTCCAGATCGAGAAACCATCTGGACCCCAACCTTCAGGCAACACTTTAAGGGTTCGGCAATCACTGTCAACCC 280

AGTTCTTCGACGGTTGATGTAAC TCAAGGTCTAGCTCTTTGGTAGGACCTGGGGTTGAAGTCCGTTGTGAATTCCTAAGCCGTTAGTGACAGTGGGG

pLM5 insert = U3

ORF U3

D E A A N Y I E S P D R E T I L O P N L O A T L . G F G N H C H P

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fig 52 pLM5 (1>5425) Site and Sequence

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CGGACAGCAGAACGCTGGCATCAGCTATCTTAGCTCCCTCCTCTCCCTCTCTCTTCAGAGCACTGGCTCTCCAGCCCCAGGAGGAGAACAGGAGGGAG
GCCGTGCTGCTTTGCGACCGTAGTCGATAGAATCGAGGAGGAGAGGGGAGAGGAGAAAGTCTCGTGACCGAGAGGTGCGGGTCTCTCTTGTCTCTCCCTC
pLM5 insert = U3
R T A E R V H Q L S . L L L S P L L F Q S T G S P A P G G E Q E G
GAGGAGATGAAGAGGAGGAGGACAGGTTCTTGGTGCTGTACCTTTGAGAACTTCCTAGGAAGGAATGGTGGGGTGGCGTTTGGGAACTTGTGCCCCCTAA
CTCTCTACTTTCTCTCTCTCTGTCGAAGAACCGACATGGAACTCTTGAAGGATCTTCTTACCACCCACCGCAAACCTTGAACACGGGGGATT
pLM5 insert = U3
G G D E R G G T G S V C C T F E N F L G R N G G V A F G N L C P L N
CACATTTACTGGCTCTCTAGAGGGCCGTTTAAACCCGCTGATCAGCTCGACTGTGCTTCTAGTTGCCAGCCATCTGTTGTTTGGCCCCCTCCCCGT
GTGTAAATGACCGAGGAGATCTCCCGGCAAAATTTGGGCGACTAGTCGGAGCTGACACGGAAGATCAACGGTCGGTAGACAACAAACGGGGAGGGGGCA
pLM5 insert = U3
T F T G L L . R A R L N P L I S L D C A F . L P A I C C L P L P R
GCCTTCTTGACCTGGAAGGTGCCACTCCCACTGCTCTTCTTAATAAAATGAGGAAATGTCATCGCATTGTCTGAGTAGGTGTCATTCTATTCTGGGG
CGGAAGGAACGGGACCTTCCACGGTGAGGGTGACAGGAAAGGATTATTTTACTCTTTAAGTAGCGTAACAGACTCATCCACAGTAAGATAAGACCCC
A F L D P G R C H S H C P F L I K . G N C I A L S E . V S F Y S G
GGTGGGGTGGGGCAGGACAGCAAGGGGGAGGATTGGGAAGACAATAGCAGGCATGCTGGGGATGCGGTGGGCTCTATGGCTTCGAGGCGGAAAGAACCA
CCACCCACCCCGCTCTGTGCTTCCCCCTCTTAACCTTCTGTTATCGTTCGTCAGACCCCTACGCCACCCGAGATACCGAAGACTCCGCCCTTTCTTGGT
G V G G A G Q Q G G G L G R O . O A C V G C G G L Y G F . G G K N Q
GCTGGGGCTTAGGGGTATCCCAACGCGCCTGTAGCGGCGCATTAAAGCGGGCGGGTGTTGGTGGTTACGCGCAGCGTGACCGCTACACTTGCCAGCGG
CGACCCGAGATCCCCATAGGGGTGCGGGGACATCGCGCGTAATTCGCGCGCCACACCAATGCGCGTCGCACTGCGCATGTGAACGGTCTCGG
L G L . G V S P R A L . R R I K R G G C G G Y A Q R D R Y T C Q R
CCTAGCGCCCGCTCTTTCTGCTTCTTCCCTTCTTCTCGCCACGTTTCGCGGCTTTCCCGCTCAAGCTCTAAATCGGGGCATCCCTTTAGGGTTCCGA
GGATCGCGGGCGAGGAAGCGAAGGAAGGAAGGAAGGAGCGGTGCAAGCGGCGGAAAGGGGAGTTCGAGATTAGCCCCGTAGGGAATCCCAGGGCT
P S A R S F R F L P F L S R H V R R L S P S S S K S G H P F R V F
TTTAGTGCTTTACGGCACCTCGACCCCAAAAACTTGATTAGGGTGATGGTTCACGTAGTGGGCCATCGCCCTGATAGACGGTTTTTCGCCCTTTGACGT
AAATCAGGAATGCCGTGGAGCTGGGGTTTTTTGAAC TAATCCCACTACCAAGTGCATCACCCTGAGCGGACTATCTGCCAAAAAGCGGAACTGCA
I . C F T A P R P Q K T . L G . V F T . V A I A L I D G F S P F D V
TGGAGTCCACGTCTTTAATAGTGGACTCTTGTTCAAAACCTGGAACAACACTCAACCTATCTCGGTCTATTCTTTTGATTATAAGGGATTTTGGGGAT
ACCTCAGGTGCAAGAAATATCACTGAGAACAAGGTTTGACCTTGTGTGAGTTGGGATAGAGCCAGATAAGAAAATAAATATTCCTTAAACCCCTA
G V H V L . . V T L V P N V N N T O P Y L G L F F . F I R D F G D
TTCCGCCATTGGTTAAAAATGAGCTGATTTAACAATAATTAACGCGAATTAATCTGTGGAATGTGTGTCAGTTAGGGTGTGGAAGTCCCCAGGC
AAGCGGATAACCAATTTTACTCGACTAAATGTTTTTAAATTCGCTTAATTAAGACACCTTACACACAGTCAATCCCAACCTTTTCAGGGTCCGA
F G L L V K K . A D L T K I . R E L I L V N V C O L G C G K S P G

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fig 52 pLM5 (1 > 5425) Site and Sequence

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CCCCAGGCAGGCAGAAGTATGCAAAGCATGCATCTCAATTAGTCAGCAACCAGGTGTGGAAGTCCCCAGGCTCCCCAGCAGGCAGAAGTATGCAAAGCA
GGGGTCCGTCCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTGGTCCACACCTTTCAGGGGTCCGAGGGGTCTCGCTTCATACGTTTCGT 3300
S P G R Q K Y A K H A S Q L V S N Q V V K V P R L P S R Q K Y A K H
TGCACTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCCATCCCGCCCTAACTCCGCCCAGTTCGCCCATTCGCCCCCATGGCTGACT
ACGTAGAGTTAATCAGTCGTGGTATCAGGGCGGGGATTGAGGCGGGTAGGGCGGGGATTGAGGCGGGTCAAGGCGGGTAAGAGGCGGGTACCGACTGA 4000
A S Q L V S N H S P A P N S A H P A P N S A Q F R P F S A P V L T
AATTTTTTTTATTATGCAAGGCGGAGGCCCTCTGCCTCTGAGCTATTCCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCTTAGGCTTTTGCAAAAAG
TTAAAAAAATAAATACGTCTCCGGCTCCGGCGGAGACGGAGACTCGATAAGGTCTTCATCACTCTCCGAAAAAACCTCCGGATCCGAAAAAGTTTTTC 4100
N F F Y L C R G R G R L C L . A I P E V V R R L F V R P R L L Q K
CTCCCGGAGCTTGATATCCATTTTCGGATCTGATCAAGAGACAGGATGAGGATCGTTTCGCATGATTGAACAAGATGGATTGCACGCAGGTTCTCCGG
GAGGGCCCTCGAACATATAGGTAAAGCCTAGACTAGTTCTCTGTCTTACTCTAGCAAGCGTACTAACTTGTTCTACCTAACGTGCGTCCAAGAGGCC 4200
A P G S L Y I H F R I . S R D R M R I V S H D . T R V I A R R F S G
CCGCTTGGGTGGAGAGGCTATTCCGGCTATGACTGGGCACAACAGACAATCCGGCTGCTCTGATGCCCGCGTGTTCGGCTGTGACGCGAGGGCGCCCGGT
GGCGAACCCACTCTCCGATAAGCCGATATGACCCGTGTTGTCTGTTAGCCGACGAGACTACGGCGGCACAAGGCCGACAGTCCGCTCCCGCGGGCCA 4300
R L G G E A I R L . L G T T D N R L L . C R R V P A V S A G A P G
TCTTTTGTCAAGACCGACCTGTCCGGTGCCCTGAATGAATGCAGGACGAGGCAGCGCGGCATCGTGGCTGGCCACGACGGGCGTTCTCTGCGCAGCT
AGAAAAACAGTTCTGGCTGGACAGGCCACGGGACTTACTTGACGTCTCTGCTCCGTGCGCCGATAGCACCGACCGGTGCTGCCCGCAAGGAACGCTCGA 4400
S F C Q D R P V R C P E . T A G R G S A A I V A G H D G R S L R S
GTGCTCGACGTTGTACTGAAGCGGGAAGGACTGGCTGCTATTGGGCGAAGTGCCGGGCGAGGATCTCCTGTCTCTCACCTTGCTCTCCGAGAAAAG
CACGAGCTGCAACAGTGACTTCGCCCTTCCCTGACCGACGATAACCCGCTTCACGGCCCCGTCTAGAGGACAGTAGAGTGGAACGAGGACGGCTCTTTC 4500
C A R R C H . S G K G L A A I G R S A G A G S P V I S P C S C R E S
TATCCATCATGGCTGATGCAATCGGGCGGCTGCATACGCTTGATCCGGCTACCTGCCCATTCGACCACCAAGCGAAACATCGCATCGAGCGAGCAGTAC
ATAGGTAGTACCGACTACGTTACGCCGCCGACGATGCGAACTAGGCCGATGGACGGTAAGCTGGTGGTTCGCTTTGTAGCGTAGCTCGCTCGTGCATG 4600
I H H G . C N A A A A Y A . S G Y L P I R P P S E T S H R A S T Y
TCGGATGGAAGCGGCTTTGTGATCAGGATGATCTGGACGAAGAGCATCAGGGGCTCGCGCCAGCCGAACGTTCGCCAGGCTCAAGGCGCGCATGCC
AGCTACCTTCGGCCAGAACAGCTAGTCTCTAGACCTGCTTCTCGTAGTCCCCGAGCGCGGTGCGGCTTGACAAGCGGTCCGAGTTCCGCGCGTACGG 4700
S D G S R S C R S G . S G R R A S G A R A S R T V R Q A Q G A H A
GACGGCGAGGATCTCGTCGTGACCATGGCGATGCCCTGCTTGCCGAATATCATGGTGGAAAAATGGCCGCTTTTCTGGATTTCATCGACTGTGGCCGCTGG
CTGCCGCTCCTAGAGCAGCACTGGGTACCGCTACGGACGAACGGCTTATAATACCTTTTACCGGCGAAAAGACCTAAGTAGCTGACACCGGCGGACCC 4800
R R R G S R R D P W R C L L A E Y H G G K V P L F V I H R L V P A G
GTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCGCTTCTCGTGTCTTACGGTAT
CACACCGCTTGGCGATAGTCTGTATCGCAACCGATGGGCACTATAACGACTTCTCGAACCGCGCTTACCCGACTGGCGAAGGAGCACGAAATGCTATG 4900
C G G P L S G H S V G Y P . Y C . R A V R R M G . P L P R A L F Y
CGCCGCTCCCGATTGCGAGCGCATCGCCTTCTATCGCCTTCTTGACGAGTTCTTCTGAGCGGGACTCTGGGGTTCGAAATGACCGACCAAGCGAGCGGCA
GGGCGAGGGCTAAGCGTCCGCTAGCGGAAGATAGCGGAAGAACTGCTCAAGAGACTCGCCCTGAGACCCCAAGCTTACTGGCTGGTTGGCTCCGGT 5000
R R S R F A A H R L L S P S . R V L L S G T L G F E M T D Q A I F

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fig 52 pLMS (1 > 5425) Site and Sequence

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ACCTGCCATCACGAGATTTCGATTCCACCGCCGCTTCTATGAAAGGTTGGGCTTCGGAATCGTTTTCCGGGACGCCGGCTGGATGATCCTCCAGCGCGS
TGGACGGTAGTGCTCTAAAGCTAAGGTGGCGGCGGAAGATACTTTCAACCCGAAGCCTTAGCAAAAGGCCCTGCGGCCGACCTACTAGGAGGTCGCGCC 510X
N L P S R D F D S T A A F Y E R L G F G I V F R D A G W M I L Q R G
GGATCTCATGCTGGAGTTCTTCGCCCACCCAAC TTGTTATTGCAGCTTATAATGGTTACAAATAAGCAATAGCATCACAAATTTACAAATAAGCA
CCTAGAGTACGACC TCAAGAAGCGGGTGGGGTTGAACAAATAACGTGGAATATTACCAATGTTTATTCGTTATCGTAGTGTTTAAAGTGTTTATTCGT 520X
D L M L E F F A H P N L F I A A Y N G Y K . S N S I T N F T N K A
TTTTTTTCACTGCATTCTAGTTGTGGTTTGTCCAAACTCATCAATGTATCTTATCATGTCTGTATACCGTCGACCTCTAGCTAGAGCTTGCGGTAATCAT
AAAAAAGTGACGTAAGATCAACACCAACAGGTTTGAGTAGTTACATAGAATAGTACAGACATATGGCAGCTGGAGATCGATCTCGAACCAGCATTAGTA 530X
F F S L H S S C G L S K L I N V S Y H V C I P S T S S . S L A . S
GGTCATAGCTGTTTCTGTGTGAAATTGTTATCCGCTCACAATTCACACAACATACGAGCCGGAAGCATAAAGTGTAAGCCTGGGGTGCCTAATGAGT
CCAGTATCGACAAAGGACACACTTTAACAATAGGCGAGTGTTAAGGTGTGTTGTATGCTCGGCCCTTCGTATTTACATTTCCGACCCACGGATTACTCA 540X
W S . L F P V . N C Y P L T I P H N I R A G S I K C K A W G A . . V
GAGCTAACTCACATTAATTGCGTTG 5425
CTCGATTGAGTGTAATTAACGCAAC
S . L T L I A L

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fig 53 pLM6 (1 > 4947) Site and Sequence

Enzymes : All 146 enzymes (No Filter)

Settings: Linear, Certain Sites Only, Standard Genetic Code

Page 1

GTGGCACTTTTCGGGAAATGTGCGCGGAACCCCTATTGTTATTTTCTAAATACATTCAAATATGTATCGGTCATGAGACAATAACCCGTGATAA-
CACCGTGAAAAGCCCCTTTACACGCGCCTTGGGGATAAACAAATAAAAGATTATGTAAGTTTATACATAGGCGAGTACTCTGTTATTGGGACTATTTA 100
V H F S G K C A R N P Y L F I F L N T F K Y V S A H E T I T L I H
GCTTCAATAATATTGAAAAGGAAGAGTATGAGTATTCAACATTTCCGTGTCGCCCTTATCCCTTTTTTTCGGGCATTTTGCTTCTGTTTTGCTCAC
CGAAGTTATTATAACTTTTTCTTCTCATACTCATAAGTTGTAAAGGCACAGCGGGAATAAGGGAAAAACGCCGTAAACGGAAGGACAAAAACGAGTG 200
A S I I L K K E E Y E Y S T F P C R P Y S L F C G I L P S C F C S
CCAGAAACGCTGGTGAAAGTAAAGATGCTGAAGATCAGTTGGGTGCACGAGTGGGTACATCGAAC TGGATCTCAACAGCGGTAAAGATCCTTGAGAGTT
GGTCTTTGCGACCACTTTCAATTTCTACGACTTCTAGTCAACCCACGTGCTCACCAATGTAGCTTGACCTAGAGTTGTCGCCATTCTAGGAACCTCAA 300
P R N A G E S K R C . R S V G C T S G L H R T G S Q Q R . D P . E F
TTCGCCCCGAAGAAGCTTTTCCAATGATGAGCACTTTTAAAGTTCTGCTATGTGGCGCGGTATTATCCCGTATTGACGCCGGCAAGAGCAACTCGGTCS
AAGCGGGGCTTTTTCGAAAAGTTACTACTCGTGAATAATTCAAGACGATACACCGGCCATAATAGGGCATAACTGCGGCCCTTCTCGTTGAGCCAGC 400
S P R R T F S N D E H F . S S A M W R G I I P Y . R R A R A T R S
CCGCATACACTATTCTCAGAATGACTTGGTTGAGTACTCACCAGTCACAGAAAAGCATCTTACGGATGGCATGACAGTAAGAGAATTATGCAGTCTGCC
GGCGTATGTGATAAGAGCTTACTGAACCAACTCATGAGTGGTCAGTGTCTTTTCGTAGAATGCCACCGTACTGTCAATCTCTTAATACGTCACGACGG 500
P H T L F S E . L G . V L T S H R K A S Y G V H D S K R I M Q C C
ATAACCATGAGTGATAACACTGCGGCCAACTTACTTCTGACAACGATCGGAGGACCGAAGGAGCTAACCCTTTTTTGCACAACATGGGGGATCATGTAA
TATTGGTACTCACTATTGTGACGCCGGTTGAATGAAGACTGTTGCTAGCCTCCTGGCTTCCTCGATTGGCGAAAAACGTTGTACCCCTTAGTACATT 600
H N H E . . H C G Q L T S D N D R R T E G A N R F F A Q H G G S C H
CTCGCCTTGATCGTTGGGAACCGGAGCTGAATGAAGCCATACCAAACGACGAGCGTGACACCACGATGCCGTAGCAATGGCAACAACGTTGCGCAAACT
GAGCGGAAC TAGCAACCTTGGCTCGACTTACTTCGGTATGGTTTGTGCTCGCACTGTGGTGTCTACGGACATCGTTACCGTTGTTGCAACGCGTTTGA 700
S P . S L G T G A E . S H T K R R A . H H D A C S N G N N V A Q T
ATTAAC TGCGAACTACTTACTCTAGCTTCCCGGCAACAATTAAGACTGGATGGAGCGGATAAAGTTGCAGGACCACCTTCGCGCTCGGCCCTTCCT
TAATTGACCGCTTGATGAATGAGATCGAAGGGCGTTGTTAATTATCTGACCTACCTCCGCCATTTTCAACGCTCTGGTGAAGACGCGAGCCGGGAAGGC 800
I N W R T T Y S S F P A T I N R L D G G G . S C R T T S A L G P S
GCTGGCTGGTTTATTGCTGATAAATCTGGAGCCGGTGAGCGTGGGTCTCGCGGTATCATTGCAGCACTGGGGCCAGATGGTAAGCCC TCCGATCGTAE
CGACCGACCAATAACGACTATTTAGACCTCGGCCACTCGCACCCAGAGCGCCATAGTAACGTCGTGACCCCGGTC TACCATTGCGGAGGGCATAGCATC 900
G V L V Y C . . I V S R . A V V S R Y H C S T G A R V . A L P Y R S
TTATCTACACGACGGGAGTCAGGCAACTATGGATGAACGAAATAGACAGATCGTGAGATAGGTGCCTCACTGATTAAGCATTTGGTAAGTCTCAGACCA
AATAGATGTGCTGCCCTCAGTCCGTTGATACCTACTTGCTTTATCTGTCTAGCGACTCTATCCACGGAGTGACTAATTCGTAACCATTTGACAGTCTGGT 1000
Y L H D G E S G N Y G . T K . T D R . D R C L T D . A L V T V R P
AGTTTACTCATATATACTTTAGATTGATTTAAACTTCAATTTTAAATTTAAAGGATCTAGGTGAAGATCCTTTTGTATAATCTCATGACCAAAATCCCT
TCAATGAGTATATGAAATCTAACTAAATTTGAAGTAAAAATTAATTTTCTAGATCCACTTCTAGGAAAACTATTAGAGTACTGTTTTAGGGA 1100
S L L I Y T L D . F K T S F L I . K D L G E D P F . . S H D O N F
TAACGTGAGTTTTCGTTCCACTGAGCGTCAGACCCGTAAGAAAGATCAAGGATCTTCTTGAGATCCTTTTTTCTGCGCGTAATCTGCTGCTTGCAAA
ATTGCACTCAAAAGCAAGGTGACTCGCAGTCTGGGGCATCTTTCTAGTTTCTTAGAAGAACCTTAGGAAAAAAGACGCGCATTAGACGACGACCTTT 1200
L T . V F V P L S V R P R R K D O R I F L R S F F S A R H L L . A H

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fig 53 pLM8 (1 > 4947) Site and Sequence

Page 2

CAAAAAACCACCGCTACCAGCGGTGGTTTGTTCGCCGATCAAGAGCTACCAACTCTTTTCCGAAGGTAAC TGGCTTCAGCAGAGCGCAGATACCAAA
GTTTTTTGGTGGCGATGGTCGCCACCAACAAACGGCCTAGTTCGATGGTTGAGAAAAAGGCTTCATTGACCGAAGTCGTCGCGTCTATGGTTT 1300
K K T T A T S G G L F A G S R A T N S F S E G N V L Q Q S A D T)
TACTGTCCTTCTAGTGTAGCGTAGTTAGGCCACCACCTCAAGAACTCTGTAGCACCGCCTACATACCTCGCTCTGCTAATCCTGTTACCAGTGGCTGCT
ATGACAGGAAGATCACATCGGCATCAATCCGGTGGTGAAGTTCTTGAGACATCGTGGCGGATGTATGGAGCGAGACGATTAGGACAATGGTCACCGACGA 1400
Y C P S S V A V V R P P L Q E L C S T A Y I P R S A N P V T S G C
GCCAGTGGCGATAAGTCGTGCTTACCAGGTTGGACTCAAGACGATAGTTACCGGATAAGGCGCAGCGGTCGGGCTGAACGGGGGTTCTGTCACACAGC
CGGTCACCGCTATTGAGCAGAAATGGCCAACTGAGTTCTGCTATCAATGGCCTATTCCGCGTCGCCAGCCGACTTGCCCCCAAGCACGTGTGTCG 1500
C Q V R . V V S Y R V G L K T I V T G . G A A V G L N G G F V H T A
CCAGCTTGAGCGAACGACCTACACGAAGTGAATACCTACAGCGTGAGCTATGAGAAAGCGCCACGCTTCCGAAGGGAGAAAGGGGACAGGTATCC
GGTCGAACCTCGCTTGCTGGATGTGGCTTGACTCTATGGATGTGCGACTCGATACTCTTTCGCGGTGCGAAGGGCTTCCTCTTTCCGCTGTCCATAGG 1600
O L G A N D L H R T E I P T A . A M R K R H A S R R E K G G Q V S
GGTAAGCGCAGGGTCGGAACAGGAGCGCACGAGGGAGCTTCCAGGGGAAACGCCTGGTATCTTTATAGTCTGTGCGGTTTCGCCACCTCTGACTT
CCATTGCGCGTCCAGCCTTGCTCTCTCGCGTGTCCCTCGAAGGTCCTCTTTGCGGACCATAGAAATATCAGGACAGCCCAAGCGGTGGAGACTGAA 1700
G K R Q G R N R R A H E G A S R G K R L V S L . S C R V S P P L T
GAGCGTCGATTTTTGTGATGCTCGTCAGGGGGGCGGAGCCTATGAAAAACGCCAGCAACGCGGCTTTTACGGTTCCTGGCCTTTTGTGCGCCTTTTG
CTCGCAGCTAAAAACACTACGAGCAGTCCCCCGCCTCGGATACCTTTTTCGGTGTGTCGCGGAAAAATGCCAAGGACCGGAAAAACGACCGGAAAAAC 1800
A S I F V M L V R G A E P M E K R Q Q R G L F T V P G L L L A F C
CTCACATGTTCTTCTCGGTTATCCCCTGATCTGTGGATAACCGTATTACCGCCTTTGAGTGAGCTGATACCGCTCGCCGAGCCGAACGACCGAGCG
GAGTGTAAGAAAGGACGCAATAGGGGACTAAGACACCATTGGCATAATGGCGGAAACCTACTCGACTATGGCGAGCGGCGTCGGCTTGCTGGCTCGC 1900
S H V L S C V I P . F C G . P Y Y R L . V S . Y R S P Q P N D R A
CAGCGAGTCAGTGAGCGAGGAAGCGGAAGAGCGCCCAATACGCAAAACGCCCTCCTCCCGCGGTTGGCCGATTCTTAATGCAGCTGGCAGCAGAGTTT
GTCGCTCAGTCAGTCGCTCCTTCGCTTCTCGCGGGTTATGCGTTTGGCGGAGAGGGGCGCAACCGCTAAGTAATTACGTCGACCGTGCTGTCCAAA 2000
O R V S E R G S G R A P N T Q T A S P R A L A D S L M O L A R Q V
CCCAGCTGGAAGCGGGCAGTGAGCGCAACGCAATTAATGTGAGTTAGCTCACTCATTAGGCACCCAGGCTTTACACTTTATGCTTCCGGCTCGTATGT
GGGCTGACCTTTCCGCCGCTCACTCGGTTGCGTTAATTACACTCAATCGAGTGAGTAATCCGTGGGGTCCGAAATGTGAAATACGAAGGCCGAGCATACA 2100
S R L E S G Q . A O R N . C E L A H S L G T P G F T L Y A S G S Y V
TGTGTGAATTGTGAGCGGATAACAATTTACACAGGAACAGCTATGACCATGATTACGCCAAGCGGCAATTAACCTCCTACATAAGGGAACAAAAGCT
ACACACCTTAACACTCGCTTATTGTTAAAGTGTCCTTTGTGATACCTGGTACTAATCGGTTTCGCGGTTAATTGGGAGTGATTTCCCTTGTTTCGA 2200
V V N C E R I T I S H R K Q L . P . L R Q A R N . P S L K G T Y A
GGGTACCGGGCCCCCTCGAGGTGACGGTATCGATAAGCTTGATATCGAATTCCTGCAGCCCGGGGATCCATGCAATAGGAGGAGGAGCCAGAGA 2300
CCCATGGCCCGGGGGGAGCTCCAGCTGCCATAGCTATTGCAACTATAGCTTAAGGACGTCGGGCCCCCTAGGTACGTTTACTCTCTCTCTCGGTCCT
G Y R A P P R G R R Y R . A . Y R I P A A R G I H A N E E E E P E

U3 stuk

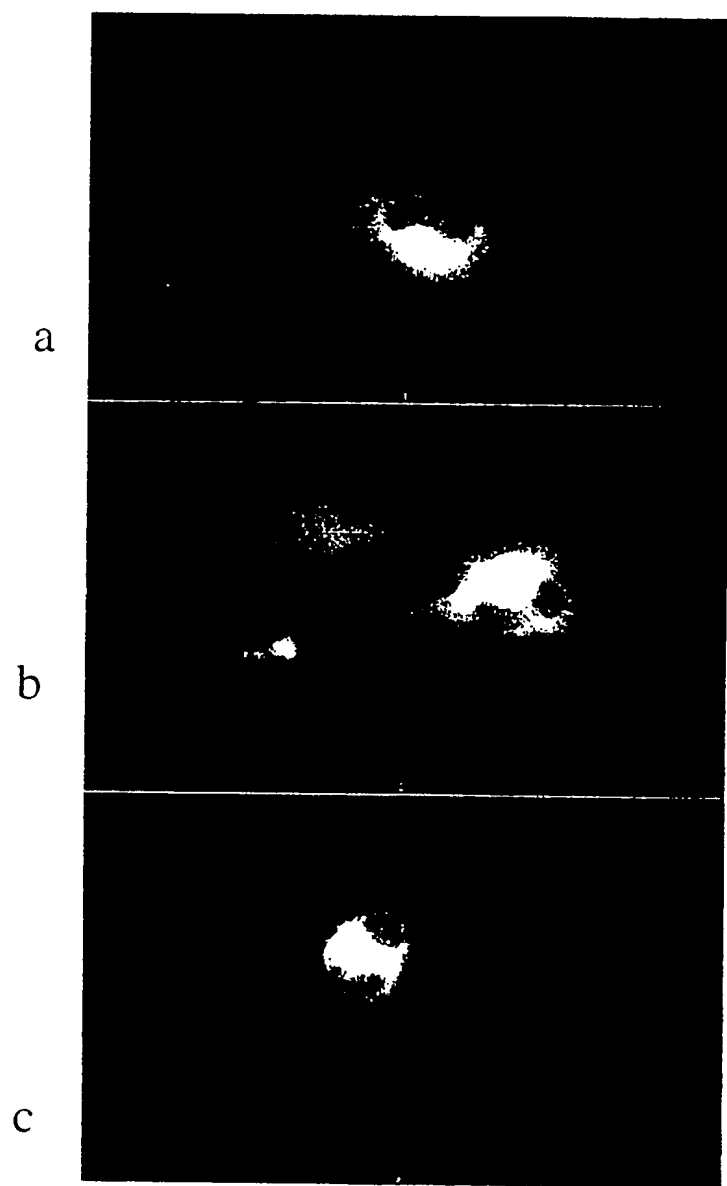


Figure 43

donderdag, 27 november 1997 15:48
fig 44 pNP9 Map (1 > 10122) Site and Sequence
Enzymes : All 148 enzymes (No Filter)
Settings : Circular, Certain Sites Only, Standard Genetic Code

Page 1

ATGACCATGATTACGCCAAGCTTGCAATGCTGCGAGGAATTCGATATCAAGCTTATCGATACCGTCGACCTCGAGGATCAGAAGAAATGGAGCAACTACC 100
CACATCCATTATGCCACCCGCGGTTTCTAAGTGAGTTAATTTTGAGTTTACGACTACAAAATGTGTTCTTTAATACTATCTTCGACTTGAGTCTATT 200
CTGTATGACTAGTTGTTGAGTGATTTTTCATTGAGAAAATATTAAAGGAACATTATTTACTTTGCTTATTTGCCCTAATTTGATTTAGTTTTCGATC 300
AACTAGATCTTACAAAATTCGAATACAAATTCATTTTCAGATTACCTCGCCACGTGTCGCCACGTGAGCAACCGCTTCAGCAACTAACCCAAATTCGA 400
ACTTTCCACAAATGTCAACATCCAGGCTTCAGACTCCACAGTCAAGATATCGAAATTTGGTAAGAATTTTATTTTGAGCTCAAACTGTATAAAATGCC 500
CAGAAAAGAGATGATAAAATGTAGTTTTCGAAAATTCACCTTTTATGCTCTAATATGACGGCTTATCTCAATTTTCTTGAGTTTATCAAA 600
AAATTTTCCACTATACAAATGTAGAAAAGTATTTTGACAAATTTGTGAGTTGACAGCTTTGTAAATAGATCCAAATGGAACCTAGATACAAGCTGTAA 700
AGTGAAGGAGCGCAAGCTATATCGGAAATAATGATCTGAAACAAATTTGTCTATTCTCAATGTTTAAAGACATGTTTGAAGATTTTCAAAATCG 800
CACTAGTTTCAGAACCTTCTTTTGTATGAAAAGTAAAAAACTATTTCAAACTCCACGCCACCATGTTTCACTCTTAATTTTATAAAATTT 900
TGCAATTTACAAATCGCTCCCTTTGTCGCGGAAAAGTCCCAAAATCAATTTCTCGGCTTCATAATGACTTTTAAATGATGTGAGAAAACACAGAAG 1000
AGGCTAACTAAATTTGACAGGGACAGGTGTGCTCTTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 1100
TTGTCATTTTCTTATAAACTTTGTGTGGAAGGAACTACACGGGAGACGGTCAATTAATTCGAATGAGAGCATGGCAATTAATCTTTTCGGAAT 1200
TGATGAATAAGATAGAGCCGATGACACTGGCTGGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAG 1300
CATCACTGACAATTAATGTGCGGTTTATGCGCTCTTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCTTCT 1400
ATTTATTTGATATTTAATTTTGTCAATTAGGATAAACGACCTTTTAAAGTTTATTTAAAAAACGATATTTTCGATTTTAAAAATCTGAAAGT 1500
TTCAAAAATCAATAAATATTCCTAACAAATGTATGGCTAAATTTTATTTCTACTGTTGACAATATCTTTATATGTATCACTGTTTTCATCTCAAA 1600
ACCTTGAATCCCCAAGTTATAGGAAGCTCCGTGTACATTTCCATGCTATGAATCGCTACTCAGCACATATCCAAAAATTAAGCTAGACGGTTGATAA 1700
TTATTGGCAGCGTAATAAAGTGCAAGCAGTTAGAATTTAATTCAGCACAGATTATCTATCAAAATCAATCTTTGAACATTCAGCCAGTTCGTACAA 1800
TTTTCCATGCTTTTGGCCATTAAAAAATTTCTCACCTCTTCATCCATCTCACTCGTATCAAAAAATATAGCAAAAGCCGACTCTACTTTTAAAG 1900
AGAAGGAGATACTGAGCCACATGGCGTGTGACCTTTTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 2000
CTTGTCTTCTTCTCGTTTGACTCGCGCTATTTTGTGGCTGCTGAAAGCCGGGAAATTTAGTATATTTAGAGCTTATCTTTATGCAATACATA 2100
AAAAACGAGGCAATTTAAAAATATTAATTAATGAGGTTGTAGATGTAGATTGGAAAAGAGAAAAAACAACAAATAGGAACCGCCAGATCAAAA 2200
TTCTATTTAAAGGTTTCAAGATGTTTAGGCAAGATTGGCTGAACAGAAAATGAAGTGCCTGCATAAATCTAGTGTAACTTTAGATTGAATCGGAA 2300
ATCCTAAGCCTGAACATATAGCTTATCTAGATCTTAGTTGCGCATAAGETCAAGCCCAAGCAGAAATGACTTGCAATTAAGTTAAGCCTAGATTGACT 2400
GCTTGCTTCAGTCTAATCCAGACTAGATTTCCAGAGAGTTTCAATTTTAAATGTTTCCAGTTTCTTGTACTTAAATCTTAATGCCCTGTGATGCGT 2500
AAAAATCGTATCCCTTCTCTCACTTTCAATTACAGATTCAATCAAGATTGGTATCAAGCCAAAGACGTCTGGACTTAAACCCCTCATCATCAACC 2600
ACTTCATCAAAATACAAATTCATTCGTCGTCAGCGCTCGAGTGGCAATAAATGTTGGCTCGACGATATCCACATCTCGCAAGAGCTTACGTA 2700
TCCGATCCTTCCGCTTCTTTTGAAGATTAATTTTCAAGATCATCATCAAGTACAGCTCTATTTGCAATCTAAACCGACCTACCTCCCACTCCA 2800
AAAACTTCTAGACCACAAACCCAGCTAGTTGCTGTTGCTACAACTACAAAAATCGGAAGCTCAAGCTAGCGCTCTGAAAGCCGTGAGCACCCCAAA 2900
CTTGCTTCTGTGAAGACTATTGGAGCAAAAAGAGCCGATAACAGCGGTGCTGCTGGTGGAAATGCTGAAATTAAGTTATTAGTAGCAAAAACC 3000

danderqg, 27 november 1997 16:48
Fig 44 pNP9 Map (1 > 10122) Site and Sequence

Page 2

CATCTTCTCATCGAATAGCCCAACCTACGAGAAAGGCGGCGGGTGCCTCAACAACTTTGTGAAAATCGCTGCCCAAGTGAAGAGTGGCT 3100
GAAGCGCGGACAGTAAGCTGGGAAGTCCACGTCTATGTGGAAGCTTTGTACGCCAAAAGTTTCTACCGTAAACGGACGCCCCAATCATATCTCAA 3200
CAAGACTCGAAACGATGCTCAAAGAGCAGTGAAGAAGAGTCCGGATACGCTGGATTCAACAGCAGCTGCCAACGTCATCATCGACGGGAAGTTCCCTAA 3300
GCATGCATTCCATCTTCCAGAGTTCAACGTCAAGCAAAAGTCTCGTCATCAGACGATCTTACTCTTAACGCCCTCCATCGTGACAGCTATCAGACA 3400
GCGGATAGCGCAACACCGGTTTCTCAAATATTATCAACAAGCTGTTGAGGAAAAACCAACTGGCAGTGAAAGGAGTGAAGACAGCGAAAAA 3500
GATCCAGCTCAGCTGTTCCGCCAGTGACACCCAGCCAAATCGGAGTTGTTAGTCCAAATTATGGCACATAAGAGTTGACAAATGACCCCGTGATAT 3600
CTGAAAAACGAGACCTGAAAGCTCCAATCAATGAGCATCGACAGCGGAGCTTCACCGCTTCCACTCTAAATCAGTTGTTCCACTTAAATGAC 3700
TTCAATCCGACAAACCAACGTACGATGTTCTTCTAAACAAGGAAAAATCACATCGCTGTCAAGTGTGTTGGATATGAGCAGTGTGCGGCTGTGAA 3800
GACTCCATTGTGGCTCATGCGTGGCTCAGGTGACTCCGCCGACAAAAATCTTGTGAATCATTGCTGGAGAGAGGATGGGAAAGAAATAGCATCAG 3900
AATCAGCGGCTACACTCTGACCGCGGTGTTGCGATGTGCGCAAAATGAGGGAGAGCTGAAAGAAATACGATGACATGACTGCTGAGCACAGAACGG 4000
CTATCTGACAACTCGAAGACAGTTCTCTTGTCTGTGGAATATCCGATAACAACGAGCTCGACGACATATCCACGGACGATTTGTCCGAGTAGAC 4100
ATGGCAACAGTGGCTCCAAACATAGCGACTATTCCTCTTGTGCGCATCCACGCTTCTTCTCTCAAGCGCCGAGTCCCGAGTGGTCTTCCACAT 4200
CAGTCGATTCTGATCTCGAGCAGAACAGGAGATGTGTACAACTTCTGTCCAGTGCCGACGACGCAACGTTGGCGCGCTGCCACTCAACTTCGG 4300
ACAACTTCGCTAAGATCCCGGGATACTCATCTATTCTCCACACTATCAGTGTGAGTGATAAGGACACATGTCTATGCACTACAGACTAGTCCA 4400
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fig 44 pNP9 Map (1 > 10122) Site and Sequence

Page 3

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fig 44 pNP9 Map (1 > 10122) Size and Sequence

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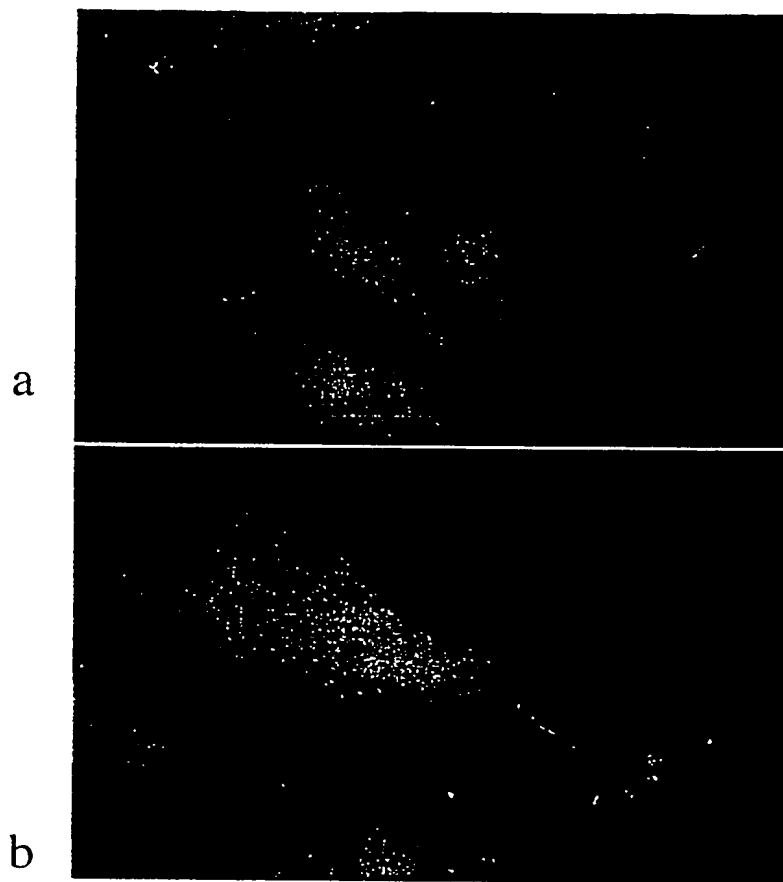


figure 45

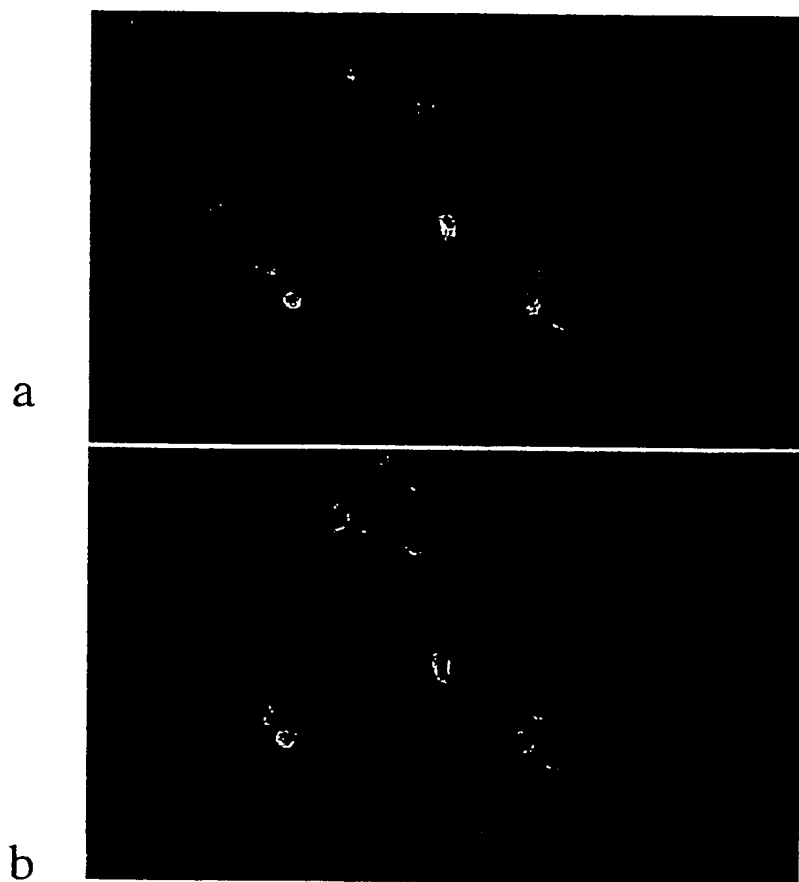


Figure 46












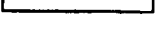
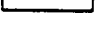
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Figure 47

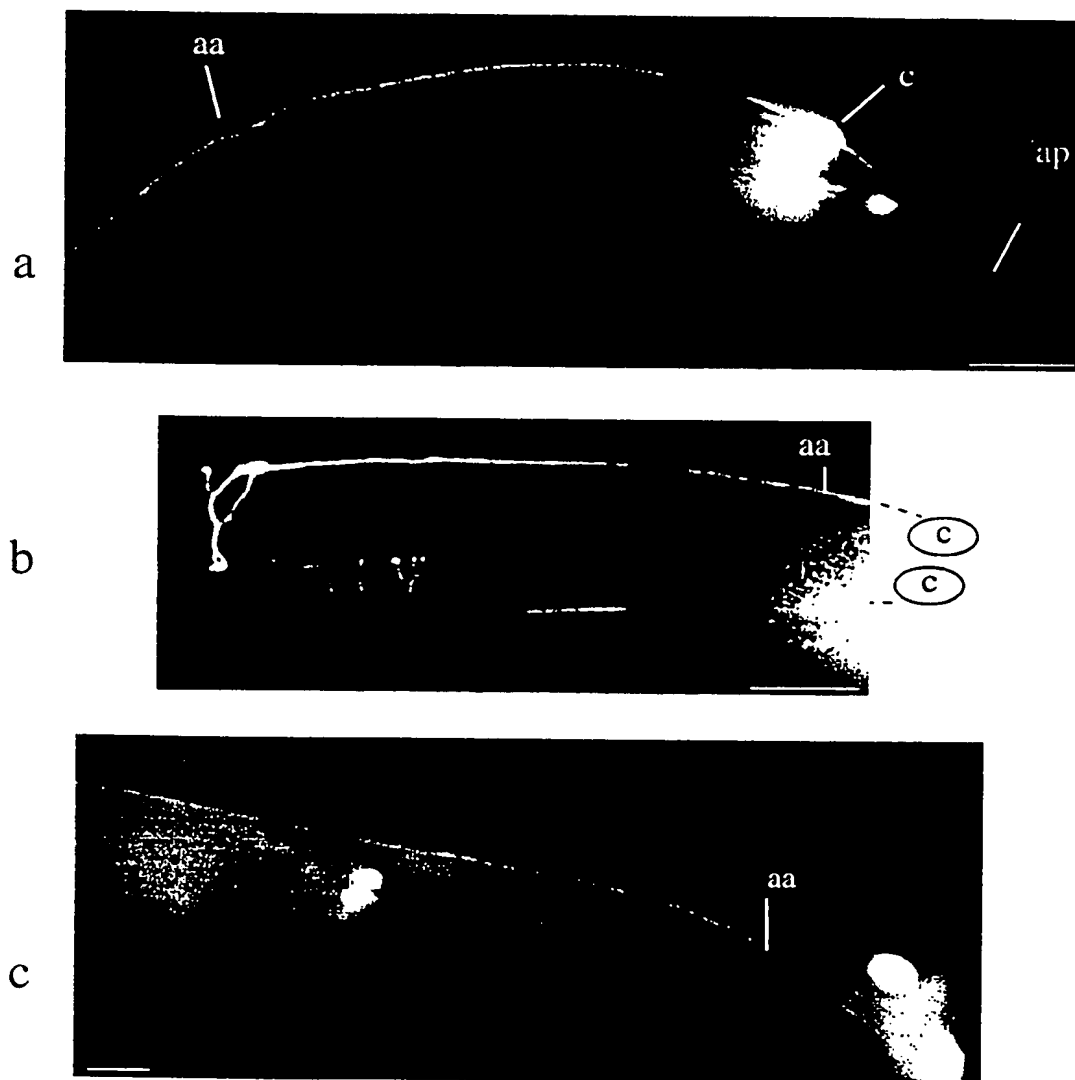


Figure 50

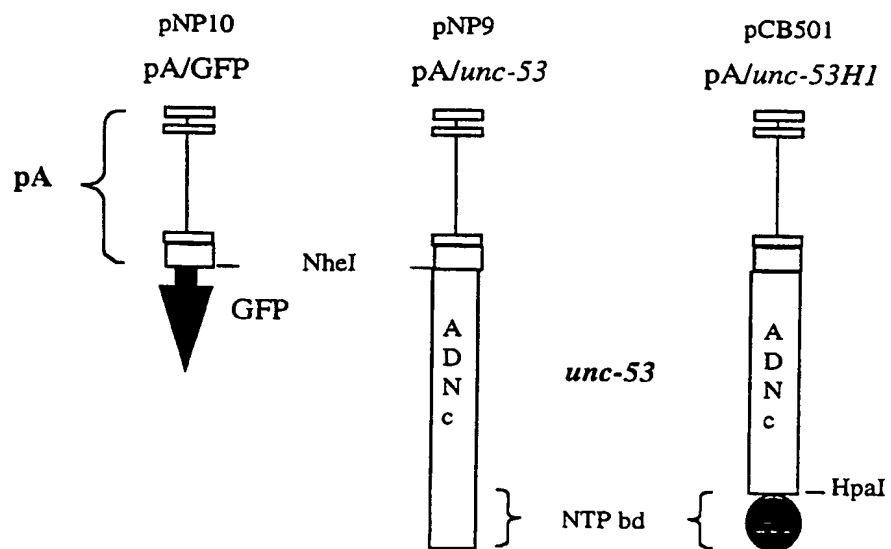


Figure 51a

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fig 35 pNP9 Map (1 > 12641) Site and Sequence

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Rg 35 pNP8 Msp (1 > 12641) Site and Sequence

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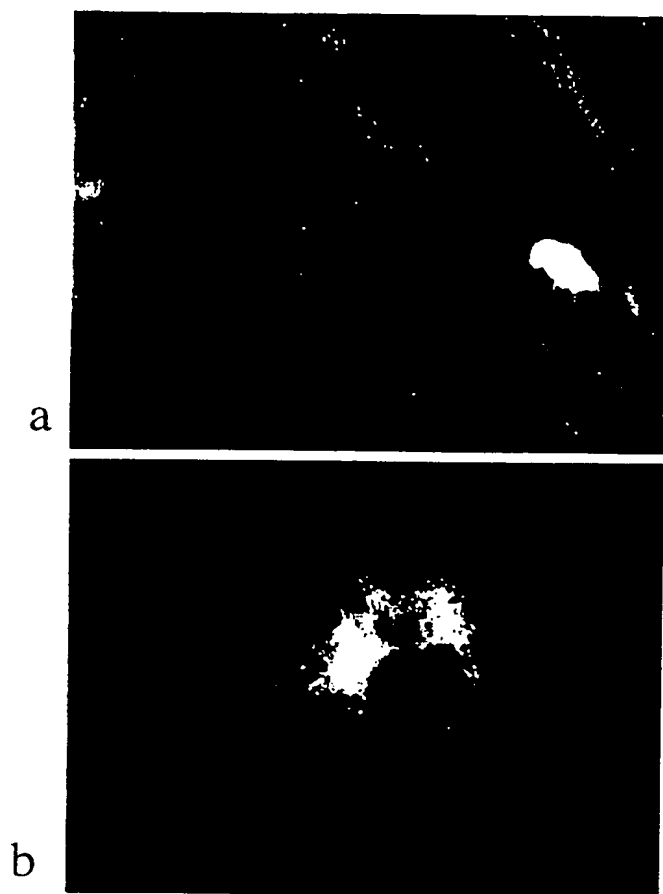


Figure 36: Association of *C. elegans* UNC-53 (expressed from pTB72) with the microtubular cytoskeleton of HepG2 cells. (A) microtubules stained with YL1/2 antibody to tubulin and (B) Staining for *C. elegans* UNC-53.

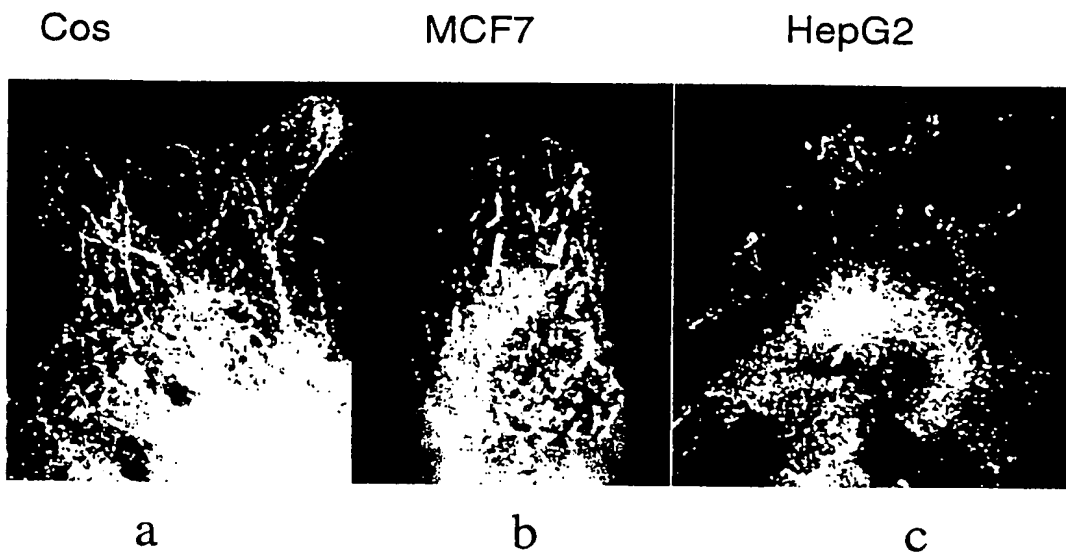


Figure 37: Microtubule (+)-end binding of *C. elegans* UNC-53 following transient transfection with pTB72 of HepG2 (a), MCF7 (b) and Cos cells (c). *C. elegans* UNC-53 was visualised by immunofluorescence using mab16-48.



Figure 38

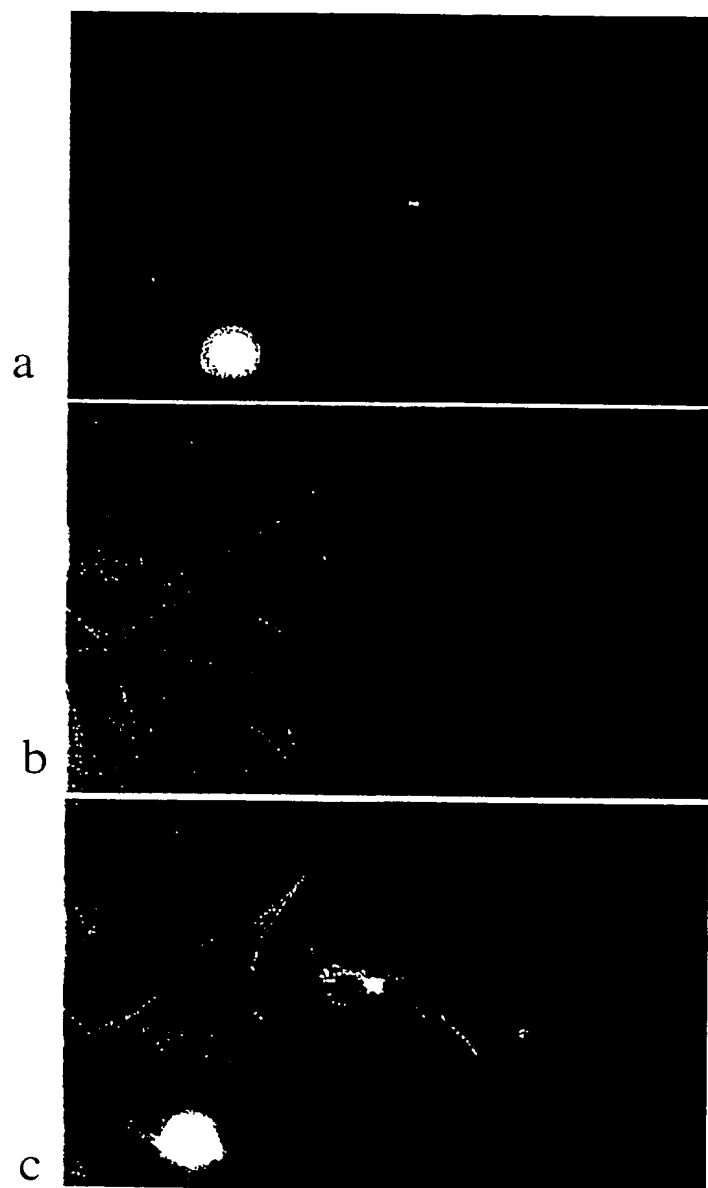


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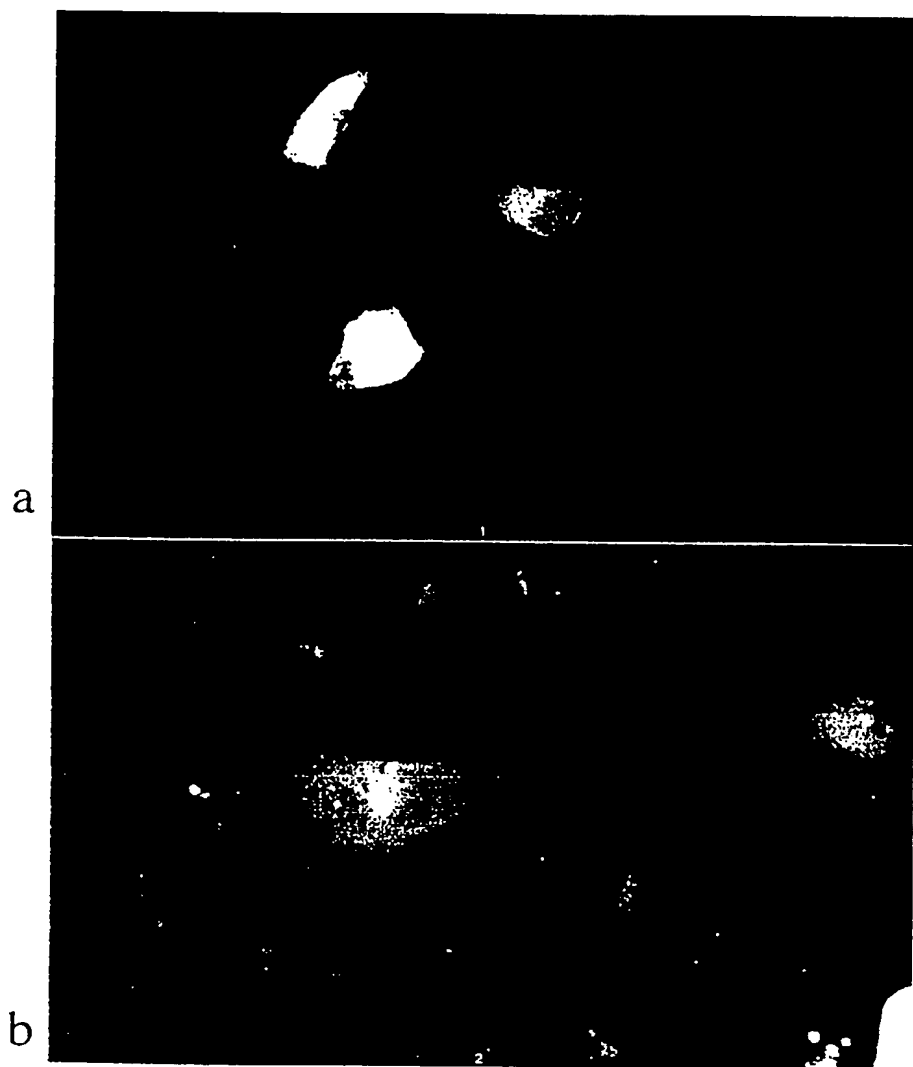


Figure 40

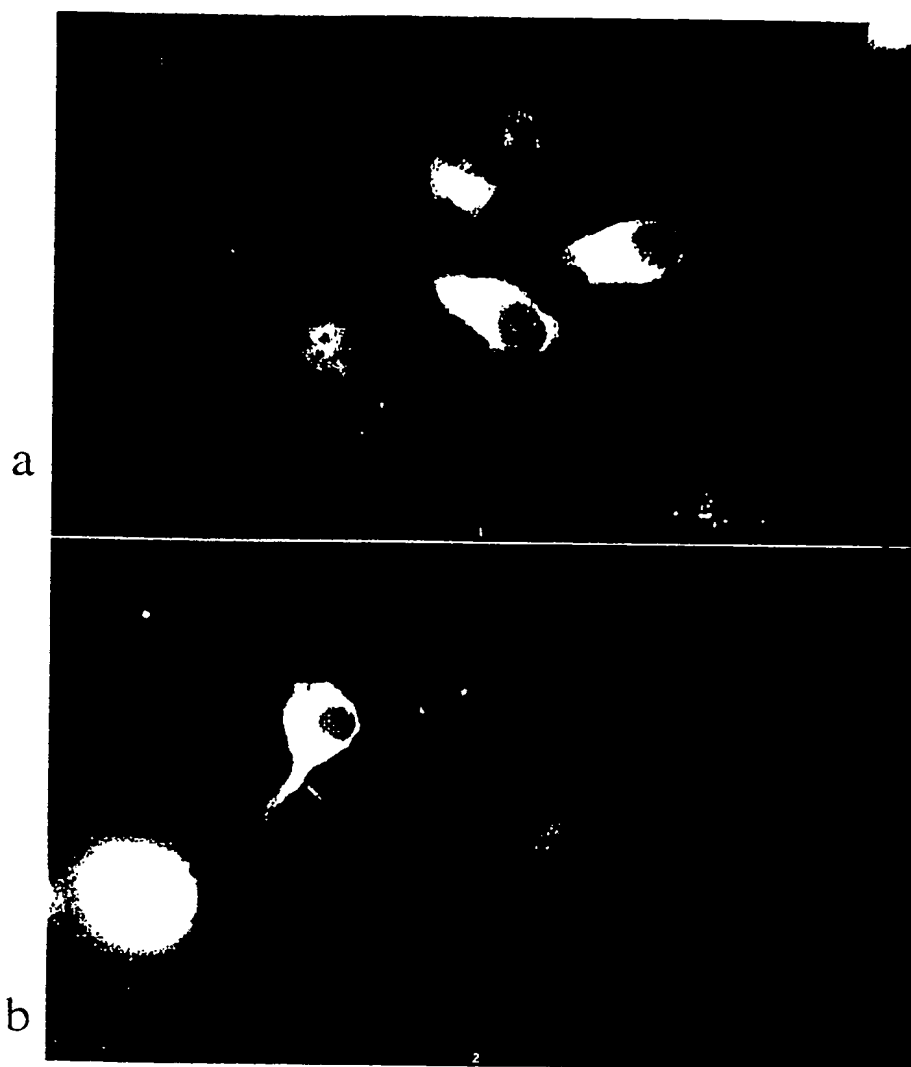


Figure 41

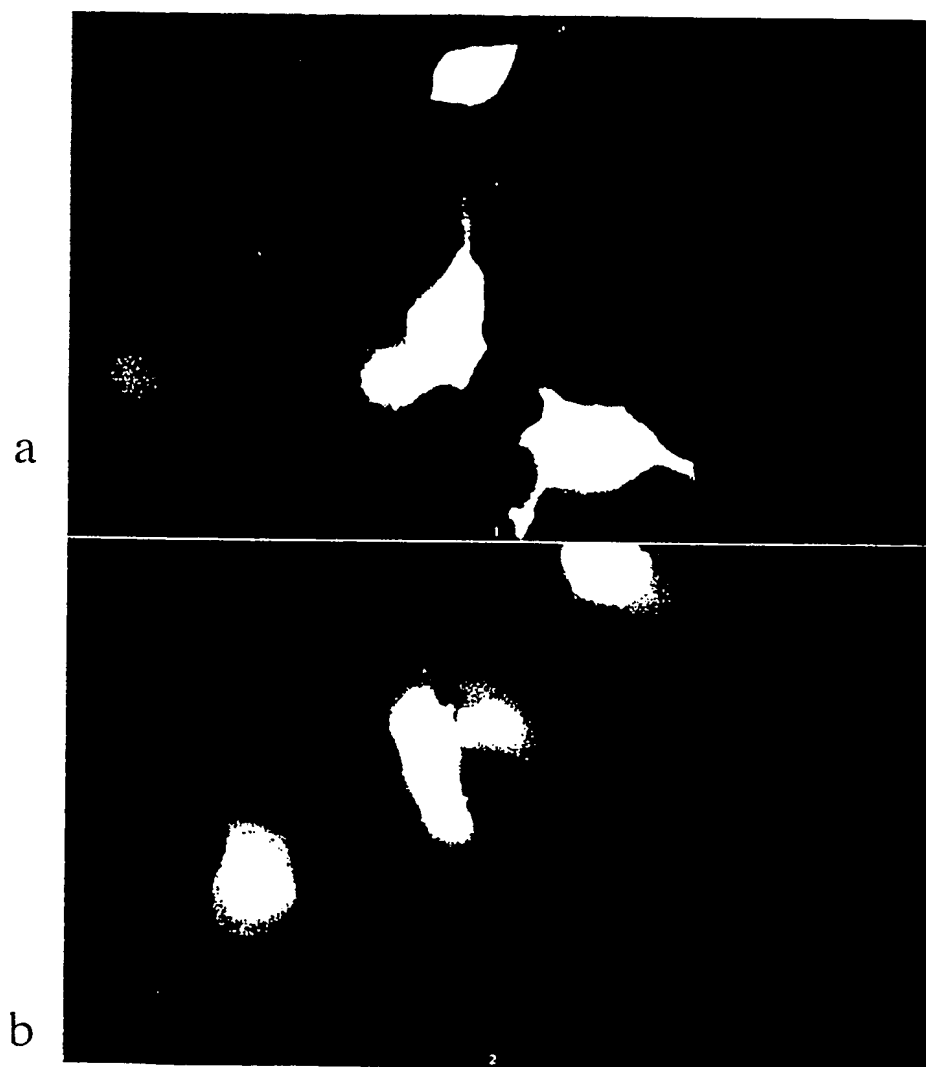


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CTACAGTGGCTCCCGGGACGAGTGGGTGCGGGGGTGTGATCCGACAAGGTACGTTTACTCTCTCTCTCGGTCTCTTCTCTCTCCATAGCCTCGACGGGA 4600

insert pLM1

ORF pLM1

C V T E G P A H P A P H T R L F H A N E E E E P E K K E V S E L R

CTGAGCTATGGGAGAAGGAAATGAAGCTTACAGACATCCGCTTGGAGGCCCTCAACTCTGCCACCAACTGGATCAGCTTCGGGAGACCATGCACAACAT
GACTCGATACCCCTCTTCTTACTTCGAATGCTGTAGGCGAACCTCCGGGAGTTGAGACGGGTGGTTGACCTAGTCGAAGCCCTCTGGTACGTGTGTA 4700

insert pLM1

ORF pLM1

S E L V E K E M K L T D I R L E A L N S A H Q L D Q L R E T I H H P

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GCAGTTGGAGGTGGACCTGCTGAAAGCAGAGAATGACCGACTGAAGGTAGCCCCAGGCCCTCATCAGGCTCCACTCCAGGGCAGGTCCCTGGATCATCT
CGTCAACCTCCACCTGGACGACTTTCGTCTCTTACTGGCTGACTTCCATC3GGGTCCGGGAGTAGTCCGAGGTGAGGTCCCGTCCAGGACCTAGTAGA

insert pLM1

ORF pLM1

O L E V D L L K A E N D R L K V A P G P S S G S T P G Q V > G S S

GCATTATCTTCCCCACGCCGCTCCCTAGGCTGGCACTCACCCATTCTTCGGCCCCAGTCTTGACAGACACAGACCTGTCACCCATGGATGGCATCAGTA
CGTAATAGAAGGGGTGCGGCGAGGGATCCGGACCGTGAGTGGGTAAAGGAAGCCGGGTCAGAACGTCGTGTCTGGACAGTGGGTACCTACCGTAGTCAT

insert pLM1

ORF pLM1

A L S S P R R S L G L A L T H S F G P S L A D T D L S P M D G I S

CTTGTGGTCCAAAGGAGGAAGTGACCTCCGGGTGGTGGTGAGGATGCCCCGACGACATCATCAAAGGGGACTTGAAGCAGCAGGAATCTTCTCGGG
GAACACCAAGTTTCTCTCTTCACTGGGAGGCCACCACTCTTACGGGGGCGTGTGTAGTAGTTTCCCTGAACTTCGTCTCTTAAGAAGGACCC

insert pLM1

ORF pLM1

T C G P K E E V T L R V V V R M P P Q H I I K G D L K Q Q E F F L G

CTGTAGCAAGGTCAAGTGGAAAAGTTGACTGGAAGATGCTGGATGAAGCTGTTTTCAAGTGTTCAGGACTATATTTCTAAATGGACCCAGCCTCTACC
GACATCGTTCCAGTCACCTTTCAACTGACCTTCTACGACCTACTTCGACAAAAGGTTCAAGTTCCTGATATAAAGATTTTACCTGGGTCGGAGATGG

insert pLM1

ORF pLM1

C S K V S G K V D V K M L D E A V F O V F K D Y I S K M D > A S T

CTGGGACTAAGCACTGAGTCCATCCATGGCTACAGCATCAGCCACGTGAAAGAGTGTGGATGCAGAGCCCCGAGATGCCTCCTTGGCGTCGAGGTG
GACCCTGATTGCTGACFCAGGTAGGTACCGATGTCGTAGTCGGTGCACCTTCTCACAACCTACGTCCTGGGGGGCTCTACGGAGGAACGGCAGCTCCAC

insert pLM1

ORF pLM1

L G L E T E S I H G Y S I S H V K R V L D A E P P E M P P C R R G

TCAATAACATATCAGTCTCCCTCAAAGGTCTGAAGGAGAAATGCGTCGACACCTCGGTGTTTCAGACGCTGATCCCCAAGCCGATGATGCAGCACTACAT
AGTTATTGTATAGTCAGAGGGAGTTTCCAGACTTCTCTTACGCAGCTGTGGACCACAAGCTCTGCGACTAGGGGTTCGGCTACTACGTCGTGTGTAT

insert pLM1

ORF pLM1

V N N I S V S L K G L K E K C V D S L V F E T L I P K P M M J H Y I

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AASCC TCCTGCTGAAGCACCGGCCTCGTCC TCTCGGGCCCCAGCGGCAGGCAAGACCTACCTGACCAATCGCTTGGCCGAGTACCTGGTGGAGCGC
TTCGGAGGACGACTTCGTGGCCGCGAGCAGGAGAGCCCGGGGTCGCCGTGCCGTTCTGGATGGACTGGTTAGCGAACCGGCTCATGEACCACCTGGCG 540
-----insert pLM1-----
-----ORF pLM1-----
S L L L K H R R L V L S G P S G T G K T Y L T N R L A E Y L V E R
TCTGGCCGTGAGGTCACAGAGGCATCGTCAGCACCTTCAACATGCACCAGCAGTCTTGCAAGGATCTGCAACTGTATCTTTCCAACCTAGCCAACCAAGA
AGACCGGCACCTCCAGTGTCTCCCGTAGCAGTCTGTGGAAGTTGTACGTGGTCTGCAGAACGTTCCTAGACGTTGACATAGAAAGGTTGGATCGGTTGGTCT 550
-----insert pLM1-----
-----ORF pLM1-----
S G R E V T E G I V S T F N M H Q Q S C K D L Q L Y L S N L A N Q
TAGACCGGGAACAGGAATGGGGATGTGCCCCCTGGTGATTCATTGGATGACCTGAGTGAAGCAGGCTCCATCAGTGAGTTGGTCAATGGGGCCCTCAC
ATCTGGCCCTTTGTCTTAACCCCTACACGGGGACCCTAAGATAACCTACTGGACTCACCCTCGTCCGAGGTAGTCACCAACCAGTTACCCCGGGAGTG 560
-----insert pLM1-----
-----ORF pLM1-----
I D R E T G I G D V P L V I L L D D L S E A G S I S E L V N G A L T
CTGCAAGTATCATAAATGTCCCTATATTATAGGTACCACCAATCAGCCTGTAAAAATGACACCAACCATGGCTTGCACTTGAGCTTCAGGATGTTGACC
GACGTTTCATAGTATTTACAGGGATATAATATCCATGGTGGTTAGTCGGACATTTTACTGTGGGTTGGTACCGAACGTGAACCTCGAAGTCTACAACCTGG 570
-----insert pLM1-----
-----ORF pLM1-----
C K Y H K C P Y I I G T T N Q P V K M T P N H G L H L S F R M L T
TCTCCAACAACGTGGAGCCAGCCAATGGCTTCCTGGTTCGTTACCTGAGGAGGAAGCTGGTAGAGTCAGACAGCGACATCAATGCCAACAAGGAAGAGC
AAGAGGTGTGTGACCTCGGTGGTTACCGAAGGACCAAGCAATGGACTCCTCCTTCGACCATCTCAGTCTGTGCTGTAGTTACGGTTGTTCCTCTCTCG 580
-----insert pLM1-----
-----ORF pLM1-----
S N N V E P A N G F L V R Y L R R K L V E S D S D I N A N K E E
TGCTTCGGGTGCTCGACTGGGTACCCAGCTGTGGTATCATCTCCACACCTTCCTTGAGAAGCACAGCACC TCAGACTTCCTCATCGGCCCTTGCTTCCT
ACGAAGCCCAAGAGCTGACCATGGGTTCGACACCATAGTAGAGGTGTGGAAGGAACCTCTCGTGTCTGGAGTCTGAAGGAGTAGCCGGGAACGAAGA 590
-----insert pLM1-----
-----ORF pLM1-----
L L R V L D V V P K L V Y H L H T F L E K H S T S D F L I G P C F F
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[illegible]

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ACAAACTCCTGGGCTTTCTGGGGAGGGGTTGAGAAAACATCAAAACACTGCAGCAGTTCCCGGAATTCAGCTTGGACTTAACAGGCTGAACCTGCTCA
TGTGAGGACCCGAAAGACCCCTCCCAAGTCTTTGTAGTTTGTGACGTCGTCAGGGGCCTTAAGTCGAACCTGAATTGGTCCGACTTGAACGAGT 570

insert pLM1

T N S V A F V G G V Q K T S K H C S S S P E F S L D L T R L N L L

AAAGAAGCCGAATTCCAGCACACTGGCGGCGTTACTAGTTCTAGATAACTGATCATAATCAGCCATACCACATTTGTAGAGGTTTACTTGCTTTAAAA
TTTCTTCGGCTTAAGGTCGTGTGACCGCGGCAATGATCAAGATCTATTGACTAGTATTAGTCGGTATGGGTAAACATCTCCAAAATGAACGAAATTTT 580

insert pLM1

K R S R I P A H W R P L L V L D N . S . S A I P H L . R F Y L L . I

AACCTCCACACCTCCCCCTGAACCTGAAACATAAAATGAATGCAATTGTTGTTAACTTGTATTGTCAGCTTATAATGGTTACAAATAAGCAATA
TTGGAGGGTGTGGAGGGGAC TTGACTTTGTATTTACTTACGTTAACAACAACAATGAACAAATAACGTCGAATATTACCAATGTTTATTTTCGTTAT 590

T S H T S P . T . N I K . M Q L L L L T C L L Q L I M V T N K A I

GCATCACAATTTTCAAAATAAGCATTTTTTTCACTGCATTCTAGTTGTTGGTTTGTCCAACTCATCAATGTATCTTAACGCGTAAATGTAAGCGTTA
CGTAGTGTTTAAAGTGTATTATTCGTAATAAAGTGACGTAAGATCAACACCAACAGGTTTGAGTAGTTACATAGAATTGCGCATTAAACATTCGCAAT 700

IT on

A S Q I S Q I K H F F H C I L V V V C P N S S M Y L N A . I V S V

ATATTTTGTAAATTCGCGTTAAATTTTGTAAATCAGCTCATTTTTTAACCAATAGGCCGAAATCGGCAAAATCCCTTATAAATCAAAAGAATAGAC
TATAAAACAATTTAAGCGCAATTTAAAAACAATTTAGTCGAGTAAAAAATGGTTATCCGGCTTTAGCCGTTTATAGGAATATTAGTTTCTTATCTG 710

IT on

N I L L K F A L N F C . I S S F F N Q . A E I G K I P Y K S K E . T

CGAGATAGGGTTGAGTGTGTTCAGTTTGAACAAGAGTCCACTATTAAAGAAGCTGGACTCCAACGTCAAAGGGCGAAAAACCGTCTATCAGGGCGAT
GCTCTATCCCACTCACAACAAGGTCAAACCTTGTCTCAGGTGATAATTTCTTGACCTGAGGTTGACGTTTCCCGCTTTTGGCAGATAGTCCCGCTA 720

IT on

E I G L S V V P V W N K S P L L K N V D S N V K G R K T V Y Q G D

GGCCCACTACGTGAACCATCACCTAATCAAGTTTTTGGGGTCGAGGTGCCGTAAAGCACTAAATCGGAACCTAAAGGGAGCCCCGATTAGAGCTT
CCGGGTGATGCACTTGGTAGTGGGATTAGTTCAAAAAACCCAGCTCCACGGCATTTCTGTGATTAGCTTGGGATTTCCTCGGGGGCTAAATCTCGAA 730

IT on

G P L R E P S P . S S F L G S R C R K A L N R N P K G S P R F R A

GACGGGAAAGCGCGCAACGTGGCGAGAAAGGAAGGAAGCAAGGAGCGGGCGCTAGGGCGCTGGCAAGTGAGCGGTACGCTGCGCGTAAC
CTGCCCCTTTCGGCCGCTTGACCGCTCTTTCTCTCCCTTCTTTGCTTTCCTCGCCCGGATCCCGGACCGTTCACATCGCCAGTGCGACGCGCATTS 740

IT on

R G K P A N V A R K E G K K A K G A G A R A L A S V A V T L R V T

CACCACCCCGCGGCTTAATGCGCCGTACAGGGCGCGTCAGGTGGCACTTTTCGGGGAATGTGCGCGGAACCCCTATTGTTTATTTTCTAAATA
GTGGTGTGGGCGGCGCAATTACGCGCGATGTCCCGCGATCCACCGTGAAGAGCCCTTTACACGCGCTTGGGGATAACAAATAAAAGATTAT 750

IT on

T T P A A L N A P L Q G A S G G T F R G N V R G T P I C L F F . I

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CATTCAAATATGTATCCGCTCATGAGACAATAACCCTGATAAATGCTTCAATAATATTGAAAAAGGAAGAGTCTGAGGCGGAAAGAACCAGCTGTGGAA
GTAAGTTTATACATAGGCGAGTACTCTGTTATTGGGACTATTTACGAAGTATTATAACTTTTCTCTTCAGGACTCCGCCCTTTCTTGGTCGACACCTT
H S N M Y P L M R Q . P . . M L Q . Y . K R K S P E A E R T S C G 760

TGTGTGTCAGTTAGGGTGTGGAAAGTCCCCAGGCTCCCCAGCAGGCAGAAGTATGCAAAGCATGCATCTCAATTAGTCAGCAACCAGGTGTGGAAAGTCC
ACACACAGTCAATCCACACCTTTTCAGGGGTCCGAGGGGTCTGCTCGCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTCCACACCTTTCAGS
M C V S . G V E S P Q A P Q Q A E V C K A C I S I S Q Q P G V E S P 770

CCAGGCTCCCCAGCAGGCAGAAGTATGCAAAGCATGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCCATCCGCCCTAACTCCGC
GGTCCGAGGGGTCTGCTCGCTTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTATCAGGGCGGGGATTGAGGCGGGTAGGGCGGGGATTGAGGCG
Q A P Q Q A E V C K A C I S I S Q Q P . S R P . L R P S R P . L R 780

CCAGTTCCGCCCATTCGCCCCATGGCTGACTAATTTTTTTTATTTATGCAAGGGCCGAGGCGCCTCGGCCTCTGAGCTATTCCAGAAGTAGTGAGG
GGTCAAGGCGGGTAAGAGGCGGGTACCGACTGATTAATAAATAAATACGCTCCGGCTCCGGCGGAGCCGGAGACTCGATAAGGTCTTCATCACTCC
P V P P I L R P M A D . F F L F M Q R P R P R P L S Y S R S S E 790

AGGCTTTTTTGGAGGCCTAGGCTTTTGCAAAGATCGATCAAGAGACAGGATGAGGATCGTTTCGCATGATTGAACAAGATGGATTGCACGCAGGTTCTCC
TCCGAAAAAACC TCCGGATCCGAAACGTTTCTAGCTAGTTCTCTGTCTACTCTAGCAAAGCGTACTAACTTGTTC TACCTAACGTGCGTCCAAGAGG
E A F L E A . A F A K I D Q E T G . G S F R M I E Q D G L H A G S P 800

GGCCGCTTGGGTGGAGAGGCTATTTCGGCTATGACTGGGCACAACAGACAATCGGCTGCTCTGATGCCGCCGTGTTCCGGCTGTACGCGCAGGGGCGGCCG
CCGGCGAACCACCTCTCCGATAAGCCGATACTGACCCGTGTTGTCGTTAGCCGACGAGACTACGGCGGCACAAAGCCGACAGTCGCGTCCCGCGGGC
A A V V E R L F G Y D V A Q Q T I G C S D A A V F R L S A Q G R P 810

GTTCTTTTTTGAAGACCGACCTGTCCGGTGCCCTGAATGAAGTGAAGACGAGGCGCGGCTATCGTGGCTGGCCACGACGGGCTTCTTGGCGCA
CAAGAAAAACAGTTCTGGCTGGACAGGCCACGGGACTTACTTGACGTTCTGCTCCGTCGCGCCGATAGCACCGACCGGTGCTGCCCGCAAGGAACGGCTC
V L F V K T D L S G A L N E L Q D E A A R L S V L A T T G V P C A 820

CTGTGCTCGACGTTGTCTGTAAGCGGGAAGGGACTGGCTGCTATTGGGCGAAGTGCCGGGGCAGGATCTCCTGTCTCTCACCTTGCTCTGCCGAGAA
GACACGASCTGCAACAGTGACTTCGCCCTTCCTTGACCGACGATAACCCGCTTCACGGCCCGTCTAGAGGACAGTAGAGTGAACGAGGACGGCTCTT
A V L D V V T E A G R D V L L L G E V P G Q D L L S S H L A P A E I 830

AGTATCCATCATGGCTGATGCAATGCGGGCGGTGCATACGCTTGATCCGGCTACCTGCCCCATTCGACCACCAAGCGAAACATCGCATCGAGCGAGCAGT
TCATAGGTAGTACCGACTACGTTACGCCGCCGACGTATGCGAATAGGCCGATGGACGGGTAAGCTGGTGGTTTCGCTTTGTAGCGTAGCTCGCTCGTGCA
V S I M A D A M R R L H T L D P A T C P F D H Q A K H R I E R A R 840

ACTCGGATGGAAGCCGGTCTTGTGATCAGGATGATCTGGACGAAGAGCATCAGGGGCTCGCGCCAGCCGAAC TGTTCGCCAGGC TCAAGGCGAGCATGC
TGAGCCTACCTTCGGCCAGAACAGCTAGTCTCTAC TAGACCTGCTCTCGTAGTCCCGAGCGGGTGGCTTGACAAGCGGTCCGAGTTCCGCTCGTACG
T R M E A G L V D O D D L D E E H Q G L A P A E L F A R L K A S H 850

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CCGACGGCGAGGATCTCGTCGTGACCCATGGCGATGCCTGCTTGCCGAATATCATGGTGGAAAAATGGCCGCTTTTCTGGATTATCGACTGTGGCCGGC- 860
GGCTGCCGCTCCTAGAGCAGCACTGGGTACCGCTACGGACGAACGGCTTAGTAGTACCACCTTTTACCGCGAAAGACCTAAGTAGCTGACACCGGCCGA
Kan/Neo
P D G E D L V V T H G D A C L P N I M V E N G R F S G F I D C G R L
GGGTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACCGTGATATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCGCTTCTCGTGCTTTACGG- 870
CCCACACCGCTGGCGATAGTCTGTATCGCAACCGATGGGCACTATAACGACTTCTCGAACCGCGCTTACCGGACTGGCGAAGGAGCACGAAATGCCA
Kan/Neo
G V A D R Y Q D I A L A T R D I A E E L G G E W A D R F L V L Y G
ATCGCCGCTCCCGATTGCGAGCGCATCGCTTCTATCGCCTTCTTGACGAGTCTTCTGAGCGGGACTCTGGGGTTCGAAATGACCGACCAAGCGACGCC 880
TAGCGGCGAGGGCTAAGCGTCCGTAGCGGAAGATAGCGGAAGAACTGCTCAAGAAGACTCGCCCTGAGACCCCAAGCTTTACTGGCTGGTTCGCTGGC
Kan/Neo
I A A P D S Q R I A F Y R L L D E F F . A G L V G S K . P T K R R
CAACCTGCCATCAGGAGATTCGATTCCACCGCCGCTTCTATGAAGGTTGGGCTTCGGAATCGTTTTCCGGGACCGCGCTGGATGATCTCCAGCGC 890
GTTGGACGGTAGTGCTCAAAGCTAAGGTGGCGGCGGAAGATATTTCAACCCGAAGCCTTAGCAAAAGGCCCTGCGGCCGACCTACTAGGAGGTCCGC
P T C H H E I S I P P P S M K G V A S E S F S G T P A G . S S S A
GGGGATCTCATGCTGGAGTTCTTCGCCACCTAGGGGGAGGCTAACTGAAACACGGAAGGAGACAATACCGGAAGGAACCCGCGCTATGACGGCAATAA 900
CCCCTAGAGTACGACCTCAAGAAGCGGGTGGGATCCCCCTCCGATTGACTTTGTGCTTCTCTGTTATGGCTTCTTGGGCGCGATATGCCGTTATT
G I S C V S S S P T L G G G . L K H G R R Q Y R K E P A L . R Q .
AAAGACAGAATAAAACGACGGTGTGGGTCGTTTGTTCATAAACCGGGGTTGGTCCCAGGGCTGGCACTCTGTGATACCCACCGAGACCCCATTC 910
TTTCTGTCTTATTTTGGTGGCACAACCCAGCAACAAGTATTTGCGCCCCAAGCCAGGGTCCCAGCGTGAGACAGCTATGGGGTGGCTCTGGGGTAAC
K D R I K R T V L G R L F I N A G F G P R A G T L S I P H R D P I
GGGCCAATACGCCCGGTTCTTCTTTTCCCCACCCACCCCAAGTTCCGGTGAAGGCCAGGGCTCGCAGCCAACGTGGGGGCGGAGGCCCTGCC 920
CCCGGTTATCGGGCGCAAGAAGGAAAGGGTGGGGTGGGGGTTCAAGCCCACTTCCGGGTCGAGCGTGGGTTGCAGCCCCGCGCTCCGGGACG
G A N T P A F L P F P P P T P Q V R V K A Q G S Q P T S G R Q A L P
ATAGCCTCAGGTACTCATATATACCTTAGATTGATTTAAACTTCATTTTAAATTTAAAGGATCTAGGTGAAGATCCTTTTGATAATCTCATGACC 930
TATCGGAGTCCAATGAGTATATGAAATCTAACTAAATTTGAAGTAAATTTAAATTTTCTAGATCCACTTCTAGGAAAACTATTAGAGTACTGGT
P Q V T H I Y F R L I . N F I F N L K G S R . R S F L I I S . P
AAATCCCTTAACGTGAGTTTTCGTTCCACTGAGCGTCAGACCCGTAGAAAAGATCAAGGATCTTCTTGAGATCCTTTTTTCTGCGCGTAATCTGCT 940
TTTAGGGAATTGCACTCAAAAGCAAGGTGACTCAGCTGCGGCATCTTTCTAGTTTCTAGAAGAACTCTAGGAAAAAGACGCGCATTAGACGAC
pUC ori
K S L N V S F R S T E R O T P . K R S K D L L E I L F F C A . S A
CTTGCAACAAAAAACCCGCTACCGCGGTGGTTTGTGGCGGATCAAGAGCTACCAACTCTTTTTCCGAAGGTAAGTGGCTTCAGCAGAGCGCA 950
GAACGTTTGTTTTTTGGTGGCGATGGTCCGACCAACAAACGCGCTAGTTCGATGGTTGAGAAAAAGGCTTCCATTGACCGAAGTCGCTCGCGT
pUC ori
A C K Q K N H R Y Q R V F V C R I K S Y Q L F F R R . L A S A E R R

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• ATACCAAACTACTGTCCTTCTAGTGTAGCCGTAGTTAGGCCACCACTTCAAGAACTCTGTAGCACCGCCTACATACCTCGCTCTGCTAATCCTGTTACCA3
TATGGTTTATGACAGGAAGATCACATCGGCATCAATCCGGTGGTGAAGTTCTTGAGACATCGTGGCGGATGTATGGAGCGAGACGATTAGGACAATGGT 960
pUC ori
Y Q I L S F . C S R S . A T T S R T L . H R L H T S L C . S C Y Q
TGGCTGCTGCCAGTGGCGATAAGTCGTGCTTACCGGGTTGGACTCAAGACGATAGTTACCGGATAAGGCGCAGCGGTGGGCTGAACGGGGGTTTCGTG
ACCGACGACGGTCACCGCTATTGACACAGAATGGCCCAACCTGAGTTCTGCTATCAATGGCCTATTCCGCGTCGCCAGCCCGACTTGCCCCCAAGCAC 970
pUC ori
V L L P V A I S R V L P G V T Q D D S Y R I R R S G R A E R G V R
CACACAGCCCAGCTTGGAGCGAACGACCTACACCGAACTGAGATACCTACAGCGTGAGCTATGAGAAAGCGCCACGCTTCCGAAGGGAGAAAGCGGGAC
GTGTGTCGGGTGCAACCTCGCTTGGTGGATGTGGCTTGACTCTATGGATGTGCACTCGATACTCTTTCGCGGTGCGAAGGGCTTCCTCTTTCGCCCTG 980
pUC ori
A H S P A V S E R P T P N . D T Y S V S Y E K A P R F P K G E R R T
AGGTATCCGGTAAGCGGCAGGGTCGGAACAGGAGAGCGCACGAGGGAGCTTCCAGGGGGAACGCCCTGGTATCTTTATAGTCCGTGTCGGGTTTCGCCACC
TCCATAGGCCATTGCGCGTCCCGAGCTTGTCTCTCGCGTGTCTCTCGAAGGTCCCCCTTTGCGGACCATAGAAATATCAGGACAGCCCAAAGCGGTGG 990
pUC ori
G I R . A A G S E Q E S A R G S F Q G E T P G I F I V L S G F A T
TCTGACTTGAGCGTCGATTTTGTGATGCTCGTCAGGGGGGGGAGCCTATGGAAAAACGCCAGCAACGCGGCCTTTTACGGTTCTTGCCCTTTTGCTG
AGACTGAACTCGCAGCTAAAAACACTACGAGCAGTCCCCCGCCCTCGGATACCTTTTTCGGTCTGTTGCGCGGAAAAATGCCAAGGACCGGAAAAACGAC 1000
pUC ori
S D L S V D F C D A R Q G G G A Y G K T P A T R P F Y G S V P F A
GCCTTTTGCTCACATGTTCTTTCTGCGTTATCCCTGATTCTGTGGATAACCGTATTACCGCCATGCAT
CGGAAAACGAGTGTACAAGAAAGGACGCAATAGGGGACTAAGACACCTATTGGCATAATGGCGGTACGTA 10070
G L L L T C S F L R Y P L I L V I T V L P P C I

donderdag, 27 november 1997 16:48
fig 35 pNP8 Map (1 > 12841) Size and Sequence
Enzymes: All 146 enzymes (No Filter)
Settings: Circular, Certain Sites Only, Standard Genetic Code

Page 1

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CACATCCATTATGCCACCCGCGGTTTCTAAGTGAGTTAATTTTGAGTTTACCACTACAAAATGTGTTCTTTAATAACTATCTTCGACTTGAGTCTATT 200
CTGTATGACTAGTTGTTGAGTGATTTTTCATTGAGAAAATATTAAGGAACATTATTTACTTTCCTTATTTGCCCTAACTTGAATTTAGTTTTTCGATC 300
AACTAGATCTTCAAAAACCTTGCAATACAATTCATTTTCAGATTACCTCGCCACGTGTCGCCACGTGAGCAACCGCTTCAGCAACTAACCCAAATTC 400
ACTTTCCCAAAATGTCAACATCCAGGCTTCAGACTCCACAGTCAAGAAATATCGAAAATTTGGTAAGAATTTTATTTTGAGCTCAAACTGTATAAAATGCC 500
CAGAAAAGAAGATGATAAAATGTAGTTTTTTTGCAAACTTCCACCTTTATGCTCTAATATGACGGCTTATATCTCAATTTTCTTGAGTTTTATCAA 600
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CATCACTGACAATTAATGTCTGGGTTTTATCGCTCTTCTCTATTCGGCACTATTCTGGGTACCAAACTGGAATACATTTTACTACTATTCAAGCC 1400
ATTTATTTTGATATTTAATTTTGTGCAATTAGGGATAAACAGGACTTTTAAAGTTTATTTAAAAAACGATATTTTCGATTTTAAAAATCTGAAAAGT 1500
TTCAAAAATCAATAAATATTCCTAACAAATGTATGGCTAAATTTTATTTCTACTGTTGACAATATCTTATATGTATCACTGTTTCCATCTCAAA 1600
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ACTTCATCAAAATACAAAATTCATTCCGTCGCTGAGCGGTTGAGTGGCAATAAATGTTGCTGACCATATCCACATCTGCGAAGAGCTTAGGTA 2700
TCCGATCTTCCGGCTCTTTTATAGAAATATATTTTCAAGATCATCATCAAGCTACAGCTCTATTTTGAATCTAAACCGACCTACCTCCCACTCCA 2800
AAAACTTCTAGACCAAAACCCAGCTAGTTGCTGTGCTACAACTACAAAATCGGAAGCTCAAGCTAGCCGCTCCGAAGCCGTGAGCACCCCAAAA 2900
CTTGCTTGTGAGACTATTGGAGCAAAACAAGACCCGATAACAGCGGTGCTGCTGCTGCTGGAATGCTGAAATTAAGTTATTCAGTAGCAAAAAC 3000

donderdag, 27 november 1997 18:46
In 35 pNFB M30 (1 > 12641) Site and Sequence

Page 2

CATCTTCCTCATCGAATAGCCCAACCTACGAGAAAGCGCGCGCTCAACAACAACTTTGTGAAATCGCTGCCCAAGTGAAGTGGCCT 3100
GAAGCGCGGACCAGTAAGCTGGGAAGTGCCACGTCTATGTGGAAGCTTTGTACGGTGAGTATTTGAAATCGGAAATGGGAATGTATTTTAAAACT 3200
GAAATTTCTACAAATAAATAAATAAAGATTTTTCTCTGATAGTATTGCATCCACTATTTTACTTTGAAGATTTATATCTTGTTTCATATTGAAG 3300
ATATCAGATATAGAAAAGAAATAAATAATTTTGACAGTTGATAATTTTGTATAGGACCAAGACAAGTGAGATATAAGCTGTCAAAGTTGATTTTC 3400
AAGAAATTTTAAACCTAGTTTTGCGAAGCTCTGGGCTCATCTATTTAGAACCGATTCGTAATCTTCCGTTCTTGACTCTACCAAAACCAAAA 3500
CCAACCTACTAATAAAATGATGAGACAATGGGAATTGTCTCCATTTTCTCTTCTCTTGACACTCTTCAGAACTATGTCCCATCTTTTGTGCT 3600
TTGTGTCCTCCCATAAAGACTCTTCCGGAATAATGTTGCAACGGAAGTGATAATTCGAGCATTTTTCGACGTGAGGGCCGAAAAACACATCTGGCTGA 3700
CAAGAGTAAAGCAATTTCTCAGCTCTTCTTCCGCTTTTCAATCTGTTTTCAAAATGAGCTACTACAGAGTGAAAGAGCACAAATTGCAAAACATT 3800
TTTGTGTGAGATGCACTTTTGAATAATTAATCTTACGTTTTAGTTTCTAGTATTTATTTTTCATATAAATTAGAGCTTCTTAGACCTCTATATTT 3900
TTTAAACTTCTACTGAAATATACGAGATTCTTTGACTTTCGGAATTGTCTTATGGCTCTATTATTTATGAGAAAACATTTTAAAAATTTTTT 4000
GAAAAAAACTGTGCATCTTCTTTTTTACATAGTAATTTCCAGCCAAAAGTTTCTACCGTAAACGGACGCCCAATCATATCTCAACAAGACTCSA 4100
AACGATGCTCAAGAGCAGTGAAGAAGAGTCCGGATACGCTGGATTCAACAGCAGCTGCCAACGTCATCATCGACGGAAGGTTCCCTAAGCATGCATTC 4200
CACATCTTCCAAGGTTCTGTTTATAGGAGAACTGTTTTTTTTGTTTCTGACCTTACATAGTCTCGGATGTTTATAAAGTGAGGTCTCTGGGACAC 4300
CTGCCATAAATGTGAATCCGCCATTTGTTGTTACAAAAACTTTTGACAGCACCTGCTTATACATTTTATGGATAAATGTCATACGGTATTTGTCAA 4400
ACCCAACTTTTAAATTTTATTTTACAGTCAAAAATGATGTTAAAGTTTAAAGATTTTACGAAAAATGTTTTACTTAAACTTTTTATATCGATA 4500
AAATTTTGAACATTCAGATAGGAGTTCCGTCCCTAAAACCTTTTGTGTCGCGGAGAACTTTTGAATTAATCTCATTTTATAAGTTCGAACTAAT 4600
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AAAGCATTGATCAAGCATGGGCTAACTGGGCGGCTGAACCATGTACATATCTTTGGCCGAGTAGTTGCAATCTAAAGATTGGAAGCTGGCTTCA 5000
AAGTCGGACTAGGCAAAAGTGCAAAAATGGAATAATCTTGAATTAATACGCTTTCCGTCTTCCATCTTCTTTTTTGTCTCTTTTGTGCGAGAT 5100
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AGTAATTTTGGATTATTTAAAGTTTGTGCGGAATAGTAAGGAGGAGTACAATTTATTTATTTGTAGAAGCTTCACTAAACTTTGATTTTCTGACCA 5500
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fig 33 pEGFPxba (1 > 5447) Site and Sequence

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AAAAGATCAGAAGAAATTGGAGCAACTACCCACATCCATTATGCCACCCGCGGTTTCTAAATTACCCTCGCCACGTGTCGCCACGTGAGCAACCGCTTCA
TTTTC TAGTCTTCTTTAACCTCGTTGATGGGTGTAGGTAATACGGTGGGCGCCAAAGATTTAATGGGAGCGGTGCACAGCGGTGACGTGCTTGGCGAAG 120

eGFPC.e.unc53xba

C.e.unc53 xba

K D Q K K L E Q L P T S I M P P A V S K L P S P R V A T S A T A S

GCAACTAACCCAAATTCCAACTTTCCACAAATGTCAACATCCAGGCTTCAGACTCCACAGTCAAGAATATCGAAAATTGATTCATCAAAGATTGGTATCA 180
CGTTGATTGGGTTTAAGGTTGAAAGGTGTTTACAGTTGTAGGTCCGAAGTCTGAGGTGTGAGTCTTATAGCTTTTAACTAAGTAGTTTCTAACCATAGT

eGFPC.e.unc53xba

C.e.unc53 xba

A T N P N S N F P Q M S T S R L Q T P Q S R I S K I D S S K I G I

AGCCAAAGACGTCTGGACTTAAACCACCCTCATCATCAACCACTTCATCAATAATACAAATTCATTCCGTCGCGTGGAGCCGTTTCGAGTGGCAATAATAA 200
TCGGTTTCTGCAGACCTGAATTTGGTGGGAGTAGTAGTTGGTGAAGTAGTTTATTATGTTTAAAGTAAGGCAGGCAGCTCGGCAAGCTCACCGTTATTATT

eGFPC.e.unc53xba

C.e.unc53 xba

K P K T S G L K P P S S S T T S S N N T N S F R P S S R S S G N N N

TGTTGGCTCGACGATATCCACATCTGCGAAGAGCTTAGAATCATCATCAACGTACAGCTCTATTTCGAATCTAAACCGACCTACCTCCCAACTCCAAAAA 210
ACAACCGAGCTGCTATAGGTGTAGAGCTTCTCGAATCTTAGTAGTAGTTGCATGTGAGATAAAGCTTAGATTGGCTGGATGGAGGGTTGAGGTTTTT

eGFPC.e.unc53xba

C.e.unc53 xba

V G S T I S T S A K S L E S S S T Y S S I S N L N R P T S Q L Q I

CCTTGGGATCCACCGGATCTAGATAACTGATCATAATCAGCCATACCACATTTGTAGAGGTTTTACTTGGCTTTAAAAAACCTCCACACCTCCCTCGAA 220
GGAACCTTAGGTGGCTAGATCTATTGACTAGTATTAGTCGGTATGGTGTAAACATCTCCAAAAAGAACGAAATTTTTTGGAGGGTGTGGAGGGGGACT

eGFPC.e.unc53xba

V D P P D L D N . S . S A I P H L . R F Y L L . K T S H T S P .

CCTGAACATAAAATGAATSCAATTGTTGTTGTTAACTTGTATTGTCAGCTTATAATGGTTACAAATAAAGCAATAGCATCACAAATTTACAAATAAA 230
GGACTTTGTATTTTACTTACGTTAAACAACAACAAATGAACAAATAACGTCGAATATTACCAATGTTTATTTCGTTATCGTAGTGTTTAAAGTGTATTAT

T . N I K . M O L L L L T C L L Q L I M V T N K A I A S Q I S Q I I

GCATTTTTTCTACTGCATTCTAGTTGTGGTTTGTCCAAACTCATCAATGTATCTTAAACGCGTAAATTTGTAAGCGTTAATTTTTGTTAAATTCGCGTTA 240
CGTAAAAAAGTGACGTGAAGTCAACACCAACAGGTTTGTAGTAGTTACATAGAATTGCGCATTTAATTCGCAATTATAAAACAATTTTAAAGCGCAE

H F F H C I L V V V C P N S S M Y L N A . I V S V N I L L K F A L

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AATTTTGTAAATCAGCTCATTTTAAACCAATAGGCCGAATCGGCAAAATCCCTTATAAATCAAAAGAATAGACCGAGATAGGGTTGAGTGTGTTCT
TTAAAAACAATTTAGTCGAGTAAAAAATGGTTATCCGGCTTTAGCCGTTTATGGGAATATTTAGTTTTCTTATCTGGCTCTATCCCAACTCACAACAA
N F C . I S S F F N Q . A E I G K I P Y K S K E . T E I G L S V V 150

CAGTTTGGAAACAAGAGTCCACTATTAAAGAACGTGGACTCCAACGTCAAAGGGCGAAAAACCGTCTATCAGGGCGATGGCCCACTACGTGAACCATCACC
GTCAAACCTTGTCTCAGGTGATAATTTCTTGACCTGAGGTTGAGTTTCCCGCTTTTGGCAGATAGTCCCGCTACCGGGTGATGCACTTGGTAGTGG
P V V N K S P L L K N V D S N V K G R K T V Y Q G D G P L R E P S P 250

CTAATCAAGTTTTTTGGGTCGAGGTGCCGTAAGCACTAAATCGGAACCCCTAAAGGGAGCCCCGATTAGAGCTTGACGGGAAAGCCGGCAACGTG
GATTAGTTCAAAAAACCCAGCTCCACGGCATTTCGTGATTAGCTTGGGATTCCCTCGGGGGCTAAATCTCGAACTGCCCTTTCGGCCGCTTGAC
S S F L G S R C R K A L N R N P K G S P R F R A . R G K P A N V 270

GGGAGAAAGGAAGGAAGAAAGCGAAAGGAGCGGGCGCTAGGGCGCTGGCAAGTGTAGCGGTACGCTGCGCGTAACCACCACACCCGCCGCGCTTAATG
CGCTCTTTCCCTTCCCTTCTTTCGCTTTCCTCGCCCGCGATCCCGCGACCGTTCACATCGCCAGTGCACGCGCATTTGGTGGTGTGGGCGCGCGCAATTAC
A R K E G K K A K G A G A R A L A S V A V T L R V T T T P A A L N 280

CGCCGCTACAGGGCGCGTCAGGTGGCACTTTTCGGGGAATGTGCGCGGAACCCCTATTGTATTATTTCTAAATACATTCAAAATATGTATCCGCTCAT
GCGGCGATGTCGCGCGCAGTCCACCGTGAAAAGCCCTTTACACGCGCCTTTGGGATAAACAATAAAAGATTATGTAAAGTTATACATAGGCGAGTA
A P L Q G A S G G T F R G N V R G T P I C L F F . I H S N M Y P L M 290

GAGACAATAACCCTGATAAATGCTTCAATAATATTGAAAAAGGAAGAGTCTGAGGCGGAAGAACCAGCTGTGGAATGTGTGTCAGTTAGGGTGTGGAA
CTCTGTTATTGGGACTATTACGAAGTTATTATACTTTTCTTCTCAGGACTCCGCCCTTCTTGGTGCACCTTACACACAGTCAATCCACACCTT
R Q . P . . M L Q . Y . K R K S P E A E R T S C G M C V S . G V E 300

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TCAGGGTCCGAGGGGTCGCTGCTTTCATACGTTTCTGACGTAGAGTTAATCAGTCGTTGGTCCACACCTTTCAGGGGTCGAGGGGTGCTCCGCTTC
S P Q A P Q Q A E V C K A C I S I S Q Q P G V E S P Q A P Q Q A E 310

TATGCAAAGCATGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCATCCCGCCCTAACTCCGCCAGTTCCGCCCATCTCCGCC
ATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTATCAGGGCGGGGATTGAGGCGGGTAGGGCGGGGATTGAGGCGGGTCAAGGCGGGTAAGAGGCGGG
V C K A C I S I S Q Q P . S R P . L R P S R P . L R P V P P I L R P 320

CATGGCTGACTAATTTTTTTTATTTATGACAGAGGCGGAGGCCGCTCGGCCTGAGCTATTCCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCCTAGGCT
GTACCGACTGATTAAAAAATAAATACGTCTCCGGCTCCGGCGGAGCCGAGAGCTCGATAAGGTCTTCATCACTCCTCCGAAAAAACCTCCGGATCCGA
M A D . F F L F M Q R P R P P R P L S Y S R S S E E A F L E A . A 330

TTTGCAAAGATCGATCAAGAGACAGGATGAGGATCGTTTCGCATGATTGAACAAGATGGATTGCACGAGGTTCTCCGGCCGCTTGGGTGAGAGGCTAT
AAACGTTTCTAGCTAGTTCTGTGCTTACTCTAGCAAAGCGTACTAATTGTTCTACCTAACGTGCGTCAAGAGGCGGGCAAGCCACCTCTCCGATA
F A K I D O E T G . G S F R M I E Q D G L H A G S P A A V V E R L 340

TCGGCTATGACTGGGCACAACAGACAATCGGCTGCTGATGCCCGGTGTTCCGGCTGTGACGCGAGSGGCGCCCGGTTCTTTTGTCAAGACCGACCT
AGCGGATACTGACCCGTGTGTGTTAGCCGACGAGACTACGGCGGCAAGGCGGACAGTCCGCTCCCGCGGGCAAGAAAAACAGTTCTGGCTGGG
F G Y D W A O O T I G C S D A A V F R L S A O G R P V L F V K T D L 350

GTCCGGTGCCCTGAATGAAGTGAAGACGAGGAGCGGGCTATCGTGGCTGGCCACGACGGGCGTTCTTGGCGAGCTGTGCTGACGTTGTCTACGAA
CAGGCCACGGGACTTACTTACGTTCTGCTCCGTCGCGCGGATAGCACCGACCGGTGCTGCCCGCAAGGAACGCGTGCACACGAGCTGCAACAGTGACT
S G A L N E L Q D E A A R L S V L A T T G V P C A A V L D V V T E 360

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GCGGGAAGGACTGGCTGCTATTGGGCGAAGTGCCGGGGCAGGATCTCCTGTCATCTCACCTTGCTCCTGCCGAGAAAGTATCCATCATGGCTGATSCAA
CGCCCTTCCCTGACCGACGATAACCCGCTTCACGGCCCCGTCTAGAGGACAGTAGAGTGAACGAGGACGGCTCTTTCATAGGTAGTACCGACTACGTT 370
A G R D V L L L G E V P G Q D L L S S H L A P A E K V S I M A D A
TGCGGGGGCTGCATACGCTTGATCCGGCTACCTGCCCATTCGACCACCAAGCGAAACATCGCATCGAGCGAGCAGTACTCGGATGGAAGCCGGTCTTGT 380
ACGCGCGGACGTATGCGAACTAGGCCGATGGACGGGTAAGCTGGTGGTTTCGCTTTGTAGCGTAGCTCGCTCGTGCATGAGCCTACCTTCGGCCAGAACA
M R R L H T L D P A T C P F D H Q A K H R I E R A R T R M E A G L V
CGATCAGGATGATCTGGACGAAGAGCATCAGGGGCTCGCGCCAGCCGAACGTGTCGCCAGGCTCAAGGCGAGCATGCCCGACGGCGAGGATCTCGTCGTG 390
GCTAGTCTCTACTAGACCTGCTTCTCGTAGTCCCCGAGCGGGTGGCTTGACAAGCGGTCGAGTTCCGCTCGTACGGGCTGCCGCTCTAGAGCAGCAC
D Q D D L D E E H Q G L A P A E L F A R L K A S M P D G E D L V V
ACCATGGCGATGCTGCTTGCCGAATATCATGGTGGAAAATGGCCGCTTTTCTGGATTTCATCGACTGTGGCCGGCTGGGTGTGGCGGACCGCTATCAGG 400
TGGGTACCGCTACGGACGAACGGCTTATAGTACCACCTTTTACCGGCGAAAAGACCTAAGTAGCTGACACCGGGCGACCCACACCGCTGGCGATACTCC
T H G D A C L P N I M V E N G R F S G F I D C G R L G V A D R Y Q
ACATAGCGTTGGCTACCCGTGATATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCGCTTCTCTGCTGCTTTACGGTATCGCCGCTCCCGATTGCGAGCG 410
TGATTCGCAACCGATGGGCACTATAACGACTTCTCGAACC GCCGTACCCGACTGGCGAAGGAGCAGCAATGCCATAGCGGCGAGGGCTAAGCGTCGC
D I A L A T R D I A E E L G G E V A D R F L V L Y G I A A P D S Q R
CATCGCTTCTATCGCTTCTTGACGAGTTCTTCTGAGCGGGACTCTGGGGTTCGAAATGACCGACCAAGCGACGCCCAACCTGCCATCAGGAGATTTCTG 420
GTAGCGGAAGATAGCGGAAGAACTGCTCAAGAAGACTCGCCCTGAGACCCCAAGCTTTACTGGCTGGTTCTGCTCGGGTTGGACGGTAGTGCTCTAAAGC
I A F Y R L L D E F F . A G L V G S K . P T K R R P T C H H E I S
ATTCCACCGCGCCTTCTATGAAGGTTGGGCTTCGGAATCGTTTTCCGGGACGCCGCTGGATGATCTCCAGCGCGGGATCTCATGCTGGAGTTCTT 430
TAAGGTGGCGCGGAAGATATTTCCAACCGAAGCCTTAGCAAAAGGCCCTGCGGCCGACCTACTAGGAGGTGCGGCCCTAGAGTACGACCTCAAGAA
I P P P P S M K G V A S E S F S G T P A G . S S S A G I S C V S S
CGCCACCTTAGGGGAGGCTAACTGAACACGGAAGGAGACAATACCGGAAGGAACCCGCGCTATGACGGCAATAAAAAGACAGAATAAACGCAAGGT 440
GCGGGTGGGATCCCCCTCCGATTGACTTTGTGCTTCTCTGTTATGGCTTCTTGGGCGGATACTGCCGTTATTTTCTGCTTTATTTTGGCTGCCA
S P T L G G G . L K H G R R Q Y R K E P A L . R Q . K D R I K R T V
GTTGGGTCGTTGTTTATAAACCGGGGTTCCGTTCCAGGGCTGGCACTCTGTGATACCCACCGAGACCCATTGGGGCCAATACGCCCGGCTTCTT 450
CAACCCAGCAACAAGTATTTGCGCCCCAAGCCAGGGTCCCGACCGTGAGACAGCTATGGGGTGGCTCTGGGGTAACCCCGGTTATGCGGGCGCAAGAA
L G R L F I N A G F G P R A G T L S I P H R D P I G A N T P A F L
CCTTTTCCCCACCCACCCCAAGTTCCGGGTGAAGGCCAGGGCTCGCAGCCAACGTCGGGGCGGCAAGGCCCTGCCATAGCTCAGGTACTCATATAT 460
GGAAAAGGGTGGGGTGGGGGTTCAAGCCCACTTCCGGGTCCCGAGCGTCGGTTGACGCCCGCCGTCGGGACGGTATCGGAGTCCAATGAGTATATA
P F P H P T P Q V R V K A Q G S O P T S G R Q A L P . P Q V T H I
ACTTTAGATTGATTTAAACTTCATTTTAAATTTAAAGGATCTAGGTGAAGATCCTTTTGTATAATCTCATGACCAAAATCCCTTAACGTGAGTTTCTG 470
TGAAATCTAACTAAATTTGAAGTAAAAATTAATTTTCTAGATCCACTTCTAGGAAAAACATTAGAGTACTGGTTTTAGGGAATTGCACTCAAAAGC
Y F R L I . N F I F N L K G S R . R S F L I I S . P K S L N V S F R
TTCCACTGAGCGTCAGACCCCGTAGAAAAGATCAAGGATCTTCTTGAGATCCTTTTTTCTGCGGTAATCTGCTGCTTGCAACAAAAAACCAACCGC 480
AAGGTGACTCGAGCTCTGGGGCATCTTTCTAGTTTCTAGAGAAGCTCTAGGAAAAAAGACGCGCATTAGACGACGAACGTTTGTTTTTTGGTGGG
S T E R O T P . K R S K D L L E I L F F C A . S A A C K Q K N H R

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fig 33 pEGFPxba (1 > 5447) Site and Sequence

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TACCAGCGGTGGTTTGTGTTGCCGGATCAAGAGCTACCAACTCTTTTCCGAAGGTAACGGCTTCAGCAGAGCGCAGATACCAAACTAGTCCTTCTAGT
ATGGTCGCCACCAACAACCGCCTAGTTCTCGATGGTTGAGAAAAAGGCTTCCATTGACCGAAGTCGTCTCGCGTCTATGGTTATGACAGGAAGATCA 500
V Q R V F V C R I K S Y Q L F F R R . L A S A E R R Y Q I L S F .
GTAGCGGTAGTTAGGCCACCACTTCAAGAACTCTGTAGCACC GCCTACATACCTCGCTCTGCTAATCCTGTTACCAGTGGCTGCTGCCAGTGGCGATAAG
CATCGGCATCAATCCGGTGGTGAAGTCTTGAGACATCGTGGCGGATGTATGGAGCGAGACGATTAGGACAATGGTCACCGACGACGGTCACCGCTATT 500
C S R S . A T T S R T L . H R L H T S L C . S C Y Q W L L P V A I S
TCGTGCTTACC GGGTGGACTCAAGACGATAGTTACCGGATAAGGCGCAGCGGTGGGCTGAACGGGGGTTTCGTGCACACAGCCAGCTTGGAGCGAA
AGCACAAGATGGCCCAACCTGAGTTCTGCTATCAATGGCCTATTCCGCGTCGCCAGCCGACTTGCCCCCAAGCACGTGTGTCGGGTCGAACCTCGCTT 510
R V L P G V T Q D D S Y R I R R S G R A E R G V R A H S P A V S E
CGACCTACACCGAACTGAGATACCTACAGCGTGAGCTATGAGAAAGCGCCACGCTTCCGAAGGGAGAAAGCGGACAGGTATCCGGTAAGCGGCAGGGT 520
GCTGGATGTGGCTTGACTCTATGGATGTGCACTCGATACTCTTTCGCGGTGCGAAGGGCTTCCCTCTTCCGCTGTCCATAGGCCATTGCGCGTCCCA
R P T P N . D T Y S V S Y E K A P R F P K G E R R T G I R . A A G
CGGAACAGGAGAGCGCAGGAGGAGCTTCCAGGGGGAAACGCCTGGTATCTTTATAGTCCTGTCGGGTTTCGCCACCTCTGACTTGAGCGTCGATTTTG 530
GCCTTGCTCTTCGCGTGTCCCTCGAAGGTCCCCCTTTCGCGGACCATAGAAATATCAGGACAGCCCAAAGCGGTGGAGACTGAACTCGCAGCTAAAAAC
S E Q E S A R G S F Q G E T P G I F I V L S G F A T S D L S V D F C
TGATGCTCGTCAGGGGGGCGGAGCCTATGGAACACGCCAGCAACGCGGCTTTTACGGTTCTTGCCCTTTTGTGGCTTTTGTGCACATGTTCTTTT 540
ACTACGAGCAGTCCCCCGCTCGGATACCTTTTTCGGTCTGTCGCCGGAATAATGCCAAGGACCGGAAAACGACCGGAAAACGAGTGTACAAGAAAG
D A R Q G G G A Y G K T P A T R P F Y G S V P F A G L L L T C S F
CTGCGTTATCCCTGATTCTGTGGATAACCGTATTACCGCCATGCAT 5447
GACGCAATAGGGGACTAAGACACCTATTGGCATAATGGCGGTACGTA
L R Y P L I L W I T V L P P C I

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fig 34 pLM4 (1 > 10070) Site and Sequence

Enzymes : 100 of 148 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

Page 1

16p

TAGTTATTAATAGTAATCAATTACGGGGTCATTAGTTCAAGCCCATATATGGAGTTCGCGTTACATAAATTACGGTAAATGGCCCGCTGGCTGACCG
ATCAATAATTATCATTAGTTAATGCCCCAGTAATCAAGTATCGGGTATATACCTCAAGGCGCAATGTATTGAATGCCATTTACCGGGCGGACCGACTGGC 100

PCMV

L L I V I N Y G V I S S . P I Y G V P R Y I T Y G K W P A V L T

CCCCAACGACCCCGCCCATTTGACGTCAATAATGACGTATGTTCCCATAGTAACGCCAATAGGGACTTTCATTGACGTCAATGGGTGGAGTATTTACGGT
GGGTTGCTGGGGCGGGTAAGTGCAGTTATTACTGCATACAAGGGTATCATTGCGGTTATCCCTGAAAGGTAAGTGCAGTTACCCACCTCATAAATGCCA 200

PCMV

A Q R P P P I D V N N D V C S H S N A N R D F . P L T S M G G V F T V

AAACTGCCCACTTGGCAGTACATCAAGTGTATCATATGCCAAGTACGCCCCCTATTGACGTCAATGACGGTAAATGGCCCGCTGGCATTATGCCAGTA
TTTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTCATGCGGGGATAACTGCAGTTACTGCCATTTACCGGGCGGACCGTAATACGGGTGAT 300

PCMV

N C P L G S T S S V S Y A K Y A P Y . R Q . R . M A R L A L C P V

CATGACCTTATGGGACTTTCCTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGATGCGGTTTGGCAGTACATCAATGGGCGTGGA
GTACTGGAATACCTGAAAGGATGAACCGTCATGTAGATGCATAATCAGTAGCGATAATGGTACCACACGCGCAAAACCGTCATGTAGTTACCGGCACCT 400

PCMV

H D L M G L S Y L A V H L R I S H R Y Y H G D A V L A V H Q V A W

TAGCGGTTTGACTCACGGGGATTTCAGTCTCCACCCCATTTGACGTCAATGGGAGTTTGTTTGGCACCAAAATCAACGGGACTTTCAGAAATGTCGTA
ATCGCCAACTGAGTGCCCTAAAGGTTAGAGGTGGGGTAAGTGCAGTTACCTCAACAAAACCGTGGTTTGTAGTTGCCCTGAAAGGTTTACAGCAT 500

PCMV

I A V . L T G I S K S P P H . R Q V E F V L A P K S T G L S K M S .

ACAACCTCCGCCCCATTGACGCAATGGGCGGTAGGCGGTACGGTGGGAGGTCATATAAGCAGAGCTGGTTTAGTGAACCGTCAGATCCGCTAGCGCTA
TGTTGAGCGGGGTAAGTGCCTTTACCGGCCATCCGCACATGCCACCTCCAGATATATTCGTCGACCAAAATCACTTGGCAGTCTAGGCGATCGCGAT 600

PCMV

Q L R P I D A N G R . A C T V G G L Y K Q S V F S E P S D P L A L

CCGGTCGCCACCATGGTGAGCAAGGGCGAGGAGCTGTTTACCGGGTGGTGCCCATCTGGTTCGAGCTGGACGGCGACGTAACGGCCACAAGTTCAGCG
GGCAGCGGTGGTACCACTCGTTCCCGCTCTCGACAAGTGGCCCCACCGGGTAGGACCAAGCTCGACCTGCCGCTGCATTTGCCGGTGTTCAGTCCG 700

EGFP

P V A T M V S K G E E L F T G V V P I L V E L D G D V N G H K F S

TGTCCGGCGAGGGCGAGGGCGATGCCACCTACGGCAAGCTGACCTGAAGTTTCATCTGCACCAACCGGCAAGCTGCCGTCGCCCTGGCCCAACCTCGTGAC
ACAGGCCGCTCCCGCTCCCGCTACGGTGGATGCCGTTTCGACTGGGACTTCAAGTAGAGCTGGTGGCCGTTTCGACGGGCACGGGACCGGTGGGAGCACTG 800

EGFP

V S G E G E G D A T Y G K L T L K F I C T T G K L P V P W P T L V T

CACCTGACCTACGGCGTGCAGTGCTTCAGCCGCTACCCCGACACATGAAGCAGCAGACTTCTTCAAGTCCGCCATGCCCGAAGGCTACGTCCAGGAG
GTGGGACTGGATGCCGACGTCACGAAGTCGGCGATGGGGCTGGTGTACTTCGTCGTGCTGAAGAAGTTCAGGCGGTACGGGCTTCGATGCAGGTCTCT 900

EGFP

T L T Y G V Q C F S R Y P D H M K Q H D F F K S A M P E G Y V Q E

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fig 34 pLM4 (1 > 10070) Site and Sequence

Page 1

CGCACCATCTTCTTCAAGGACGACGGCAAC TACAAGACCCGCGCCGAGGTGAAGTTCGAGGGCGACACCCCTGGTGAACCGCATCGAGCTGAAGGGCAGC
GCGTGGTGAAGAAGTTCCTGCTGCCGTTGATGTTCTGGGCGCGGCTCCACTTCAAGCTCCCCTGTGGGACCCTTGGCGTAGCTCGACTTCCCGTAGC 1000

EGFP

R T I F F K D D G N Y K T R A E V K F E G D T L V H R I E L K G I

ACTTCAAGGAGGACGGCAACATCCTGGGGCACAAGCTGGAGTACAAC TACAAGCCACAACGCTCTATATCATGGCCGACAAAGCAGAAGAACGGCATCAA
TGAAGTTCCTCCTGCCGTTGTAGGACCCCGTTCGACCTCATGTTGATGTTGTCGGTGTTCAGATATAGTACCGGCTGTTCTGCTTCTTCCCGTAGTT 1100

EGFP

D F K E D G N I L G H K L E Y N Y N S H N V Y I M A D K Q K N G I

GGTGAAC TTCAAGATCCGCCACAACATCGAGGACGGCAGCGTGCAGCTCGCCGACCAC TACCAGCAGAACACCCCATCGGCGACGGCCCGTGTCTGTG
CCACTTGAAGTTCTAGGCGGTGTGTAGCTCCTGCCGTCGCACGTCGAGCGGCTGGTGATGGTCTGCTTGTGGGGGTAGCCGCTGCCGGGGCACGACGAC 1200

EGFP

V N F K I R H N I E D G S V Q L A D H Y Q Q N T P I G D G P V L L

CCCGACAACCACTACCTGAGCACCCAGTCCGCCCTGAGCAAAGACCCCAACGAGAAGCGCGATCACATGGTCTGCTGGAGTTCGTGACCGCGCGCGGGA
GGGCTGTTGGTGATGGACTCGTGGGTGAGCGGGGACTCGTTTCTGGGGTGTCTTCTCGCGCTAGTGTACCAGGACGACCTCAAGCACTGGCGCGCGCCCT 1300

EGFP

P D N H Y L S T Q S A L S K D P N E K R D H M V L L E F V T A A G

TCACTCTCGGCATGGACGAGCTGTACAAGTCCGGACTCAGATCTCGAGCTCAAGCTTCGAATCTGCAGTCGATAAGCTTGATATCGAATTCCTGCAGCC
AGTGAGAGCCGTACCTGCTCGACATGTTCAAGGCTGAGTCTAGAGCTCGAGTTCGAAGCTTAAGACGTCAGCTATTCTGAACATAGCTTAAGGACGTCGG 1400

EGFP

I T L G M D E L Y K S G L R S R A Q A S N S A V D K L D I E F L Q P

CCTGCTCTCAGCCAGATGCTGGACCCAGAGTCCCAGAGAAAGAGGACAGTGCAGAATGTCTGGATCTCCGGCAGAACCTGGAAGAGACCATGTCCAGC
GGACGAGAAGTCGGTCTACGACCTGGGTCTCAGGGTCTCTTTCTCTGTCACGCTTACAGGACCTAGAGGCCGCTTGGACCTTCTCTGGTACAGGTGCG 1500

Insert pLM1

ORF pLM1

L L F S D M L D P E S O R K R T V Q N V L D L R Q N L E E T M S S

CTGCGAGGGTCCAGGTGACTCACAGCTCCCTGGAGATGACC TGCTACGACAGCGATGATGCCAACCCACGAGCGTGTCAGCCCTCTCAACCGCTCG
GACGCTCCAGGGTCCACTGAGTGTGAGGGGACCTCTAC TGGACGATGCTGTCGCTACTACGGTGGGTGCGTCCGACAGGTCCGAGAGGTTGGCGAGCA 1600

Insert pLM1

ORF pLM1

L R G S O V T H S S L E M T C Y D S D D A N P R S V S S L S N R S

CCCCTCTGTCATGGCGCTATGGCCAGTCCAGTCCGCGGCTGCAGGCTGGTGACGCGCCCTCTGTGGGTGGGAGCTGCCGCTCGGAGGGGACGCCCCCTG
GGGAGACAGTACCGGATACCGGTGAGGTGAGGCGCCGACGTCGACGAC TGGCGGGAGACACCCACCTTCGACGGCGAGCTCCCC TGGGGGCGGAC 1700

Insert pLM1

ORF pLM1

S P L S W R Y G O S S P R L Q A G D A P S V G G S C R S E G T P A V

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fig 34 pLM4 (1 > 10070) Site and Sequence

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GTACATGCACGGCGAACGGGCCACTAC TCCACACCATGCCATGCGCA3CCCCAGCAAGCTCAGCCATATCTCCCGCTGGAGCTGGTCGAATCCCTG
CATGTACGTGCCGCTTGCCCGGGTGATGAGGGTGTGGTACGGGTACGGCTCGGGGTCTTCGAGTCGGTATAGAGGGCGGACCTCGACCAAGCTTAGGGAC

insert pLM1

ORF pLM1

Y M H G E R A H Y S H T M P M R S P S K L S H I S R L E L V E S L

GACTCGGATGAGGTGGACCTCAAGTCCGGCTACATGAGCGACAGTGACCTCATGGGCAAGACCATGACGGAGGATGATGACATCACTACCGGCTGGGATG
CTGAGCCTACTCCACCTGGAGTTCAGGCCGATGTACTCGCTGTCTAC TGGAGTACCGCTTC TGGTACTGCCCTCC TACTACTGTAGTGTAGTGGCCGACCTAC

insert pLM1

ORF pLM1

D S D E V D L K S G Y M S D S D L M G K T M T E D D D I T T G V D

AAAGCAGCTCCATCAGTAGTGGACTCAGCGATGCCCTCAGACAATCTCAGTTCAGAAGAATCAATGCCAGCTCCTCACTCAACTCCCTCCCAAGTACTCC
TTTCGTCGAGGTAGTCATCACCTGAGTCGCTACGGAGTCTGT TAGAGTCAAGTCTCTTAAGTTACGGTCGAGGAGTGAGTTGAGGGAGGGTTTCATGAGG

insert pLM1

ORF pLM1

E S S S I S S G L S D A S D N L S S E E F N A S S S L N S L P S T P

CACTGCTTCTCGCAGGAACCAACAATAGTGCTACGCACAGACTCAGAGAAGCGCTCACTGGCAGAAAGTGGCTGAGCTGGTTTAGTGAATCAGAGGAG
GTGACGAAGAGCGTCTTGAGTTGTTATCAGGATGCGTGTCTGAGTCTCTCGCGAGTGACCGTCTTTCACCCGACTCGACCAAACTCACTTAGTCTCTTC

insert pLM1

ORF pLM1

T A S R R N S T I V L R T D S E K R S L A E S G L S V F S E S E E

AAAGCCCCATAAAAACTGGAGTACGACAGTGGTAGCC TGAAGATGGAACCTGGGACTTCTAAGTGGCGGAGGGAGCGGCTGAGAGCTGTGATGATTCAT
FTTCGGGGATTTTGTGACCTCATGCTGTCAACATCGGACTTCTACCTTGGACCCTGAAGATTACCGCCTCCCTCGCCGGACTCTCGACACTACTAAGTA

insert pLM1

ORF pLM1

K A P K K L E Y D S G S L K M E P G T S K V R R E R P E S C D D S

CCAAGGGTGGAGAAGTGA AAAAGCCATCAGCCTGGGCCACCTGGTTCCTTGAAGAAGGGCAAGACCCACC TG TGGCTGTAACTTCCCCATCACTCA
GGTTCACCTCTTGACTTTTTCGGTAGTCGGACCCGGTGGGACCAAGGGACTTCTTCCCGTTC TGGGGTGGACACCGACATTGAAGGGGGTAGTGAGT

insert pLM1

ORF pLM1

S K G G E L K K P I S L G H P G S L K K G K T P P V A V T S P I T H

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fig 34 pLM4 (1 > 10070) Site and Sequence

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CACAGCCCAGAGTGGCCCTCAAAGTCGCAGGCAAACTGAGGGCAAAGCTACAGACAAGGGTAAGCTTGCAGTGAAGAACTAGGGCTCCAAACGCTCCTCC
GTGTCGGGTCTACGGGAGTTTCAGCGTCCGTTTGACTCCCGTTTCGATGCTGTTCCTCATCGAACGTCACCTTATGACCCGAGGTTGCGAGGAGG 3400

insert pLM1

ORF pLM1

T A Q S A L K V A G K P E G K A T D K G K L A V K N T G L Q R S S

TCTGATGCTGGTGGGACCGCCTGAGTGTGCTAAGAAGCCCCCTCGGGCATTGCTCGCCCTCCACTTCGGGATCCTTCGGCTACAAGAAGCCTCCTC
AGACTACGACCAGCCCTGGCGGACTCCTACGATTCTTCGGGGGAGCCCGTAACGAGCGGGGAGGTGAAGCCCTAGGAAGCCGATGTTCTTCGAGGAGG 3500

insert pLM1

ORF pLM1

S D A G R D R L S D A K K P P S G I A R P S T S G S F G Y K K P P

CTGCCACAGGCACAGCCACTGTATGCAAACCTGGTGGTTCAGCCACTCTCAGCAAGATCCAGAAGTCCTCAGGCATCCCTGTCAAGCCAGTAAATGGGGG
GACGGTGTCGCTGCGGTGACAGTACGTTTGACCACCAAGTCGGTGAGAGTCTGTTCTAGGCTTCAGGAGTCCGTAGGGACAGTTTCGGTCATTTACCCGC 3600

insert pLM1

ORF pLM1

P A T G T A T V M Q T G G S A T L S K I Q K S S G I P V K P V N G R

CAAGACTAGCTTAGATGTTTCCAACAGCGCAGAGCCAGGATTCTTGGCTCCTGGAGCCCGTTCTAACATCCAGTACCGCAGCCTGCCCGGCCAGCCAAAG
GTTCTGATCGAATCTACAAAGGTGTCGCGTCTCGGTCTTAAGGACCGAGGACCTCGGGCAAGATTGTAGGTCATGGCGTCGGACGGGCGCGTTCGGTTC 3700

insert pLM1

ORF pLM1

K T S L D V S N S A E P G F L A P G A R S N I Q Y R S L P R P A I

TCAAGTTCATGAGCGTGACCGGCGGGGGGGTGGACCTCGCCCTGTGAGCAGCAGCATTGACCCAGTCTCCTCAGCACCAAGCAGGAGGCCCTTACGC
AGTTCAAGATACTCGCACTGGCCGCCGCCACCCTGGAGCGGGACACTCGTCTGCTAAGTGGGTCAGAGGAGTCTGGTTCTGCTCCCTCCGGAATGGG 3800

insert pLM1

ORF pLM1

S S S M S V T G G R G G P R P V S S S I D P S L L S T K Q G G L T

CTTCCAGACTGAAGGAGCCTACCAAGGTAGCCAGTGGGCGGACCCTCCAGCCCTGTCAATCAGACAGATCGGGAAAAGGAGAAGGCCAAAGCCAAGGG
GAAGGTCGACTTCTCGGATGGTTCCATCGGTACCCGCCCTGGTGAGGTCGGGACAGTATGCTGTCTAGCCCTTTTCTCTTCGGGTTTCGGTTCGG 3900

insert pLM1

ORF pLM1

P S R L K E P T K V A S G R T T P A P V N Q T D R E K E K A X A I A

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fig 34 pLM4 (1 > 10070) Site and Sequence

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AGTGGCCTTGGACTCAGACAACATCTCC TTGAAGAGTATTGGCTCCCCAGAGAGTACTCCCAAGAACCAAGCAAGCCACCCACAGCCACCAAGCTGGCA
TCACCGGAACCTGAGTCTGTTGTAGAGGAACCTTCATAACCGAGGGGTCTCTCATGAGGGTTC TTGGTTCGTTCCGGTGGGGTGTCGGTGGTTCGACCGT 300

insert pLM1

ORF pLM1

V A L D S D N I S L K S I G S P E S T P K N Q A S H P T A T K L A

GAGCTGCCACCAACCCCTCTCAGGGCCACAGCGAAGAGCTTTGTCAAACCAACCCTCACTAGCCAATCTTGACAAGGTCAACTCCAACAGTCTGGATCTAC
CTCGACGGTGGTTGGGAGAGTCCCGGTGTCGCTTCTCGAAACAGTTTGGTGGGAGTGATCGGTTAGAAGTGTTCAGTTGAGGTTGTCAGACCTAGATG 310

insert pLM1

ORF pLM1

E L P P T P L R A T A K S F V K P P S L A N L D K V N S N S L D L

CATCATCCAGTGATACCAACCATGCTTCAAAGGTCCCAGATCTGCATGCTACAAGCTCAGCATCTGGGGGCCCTCTCCCTTCTGCTTACCCCCAGTCC
GTAGTAGGTCACTATGGTGGGTACGAAGTTTCCAGGGTCTAGACGTACGATGTTGAGTTCGTAGACCCCGGGAGAGGGAAGGACGAAGTGGGGGTCAGG 320

insert pLM1

ORF pLM1

P S S S D T T H A S K V P D L H A T S S A S G G P L P S C F T P S P

GGCACCATCTCAATATTAAGTCAAGCCAGCTTCTCCCAGGGCTGGAGCTAATGAGTGGTTTCAGTGTGCCAAAAGAGACCCGATGTACCCCAAATC
CCGTGGGTAGGAGTTATAATTGAGTCGGTCGAAGAGGGTCCCGGACCTCGATTACTACCAAAGTCACACGGTTTCTCTGGGCGTACATGGGGTTTGAG 330

insert pLM1

ORF pLM1

A P I L N I N S A S F S Q G L E L M S G F S V P K E T R M Y P K L

TCAGGCCTGCACAGGAGCATGGAGTCCCTCCAGATGCCAATGAGCCTCCCAAGTGCCTTCCCCAGCAGTACTCCCGTCCCACCCACCTGCTCCCCCTG
AGTCCGGACGTGCTCTCGTACCTCAGGGAGGTCTACGGTTAC TCGGAGGGGTACGGAAGGGTTCGTCATGAGGGCAGGGGTGGGGTGGACGAGGGGGAC 340

insert pLM1

ORF pLM1

S G L H R S M E S L Q M P M S L P S A F P S S T P V P T P P A P P

CTGCTCCACAGAAGAAGAGACGGAAGAGCTGACTTGGAGTGGAAAGCCCCAGAGCTGGGCAACTGGACAGTAATCAGCGGGATCGGAACACTCTTCCCAA
GACGAGGGTGTCTTCTTCTGCTTCTCGACTGAACCTCACCTTCGGGGTCTCGACCCGTTGACCTGTGATTAGTCGCCCTAGCCTTGTGAGAAGGGTT 350

insert pLM1

ORF pLM1

A A P T E E E T E E L T V S G S P R A G O L D S N O R D R N T L P I

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fig 34 pLM4 (1 > 10070) Site and Sequence

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GAAAGGGCTCAGGTACCAGCTTCAGTCCAGGAGGAGACCAAGGAGAGGCGACATTCCCATACCATTTGGTGGGCTGCCGTAATCCGATGACCAGTCAGAG
CTTTCCGAGTCCATGGTCAAGTCAGGGTCTCTCTGTTCTCTCCGCTGTAAGGGTATGGTAACCAACCCGACGGACTTAGGCTACTGGTCAGTCTC

3600

insert pLM1

ORF pLM1

K G L R Y Q L Q S Q E E T K E R R H S H T I G G L P E S D D Q S E

CTGCCTTCTCCCTGCACCTTCCCATGTCTCTGAGTGCAAAGGGCAACTTACCAACATAGTGAGTCCACATGCGGCCACCCAGCCAAGAATCACCCGCT
GACGGAAGAGGGGGACGTGAAGGGTACAGAGACTCACGTTTCCCGGTGAATGGTGTATCACTCAGGGTGACGCCGGTGGTGGGTTCTTAGTGGGGCA

3700

insert pLM1

ORF pLM1

L P S P P A L P M S L S A K G Q L T N I V S P T A A T T P R I T R

CCAACAGCATCCCCACCCACGAGGGCGCCTTCGAGCTGTACAGCGGCTCCCAAATGGGGAGCACCCGTGTCCTGGCCGAGAGACCCAAGGGAATGATTCC
GGTTGTCGTAGGGGTGGGTGCTCCGCCGAAGCTCGACATGTGCGCCGAGGGTTTACCCCTCGTGGGACAGGGACCGGCTCTCTGGGTTCCCTTACTAAGC

3800

insert pLM1

ORF pLM1

S N S I P T H E A A F E L Y S G S Q M G S T L S L A E R P K G M I R

GTCAGGATCCTTCGAGACCCACGGACGATGTTACGGCTCAGTGCTGTCCCTGGCCTCCAGTGCCCTCTCCACCTACTCCTCAGCTGAGGAGAGGATG
CAGTCTTAGGAAGGCTCTGGGGTGCCTGCTACAAGTGCCGAGTCACGACAGGGACCGGAGGTCACGGAGGAGGTGGATGAGGAGTCGACTCCTCTCTCTAC

3900

insert pLM1

ORF pLM1

S G S F R D P T D D V H G S V L S L A S S A S S T Y S S A E E R M

CAATCTGAGCAAATCCGAAGCTTCGTAGGGAACGGAATCATCCAGGAAAAGTGCCACCTTGACGTCACAGCTTCTGCAATGCTAATCTGGTGG
GTTAGACTCGTTTAGGCCTTCGAAGCATCCCTTGACCTTAGTAGGGTCTTTTACCCGGTGAACGTCAGAGTCGAAAGACGGTTACGATTAGACCAAC

4000

insert pLM1

ORF pLM1

D S E O I R K L R R E L E S S Q E K V A T L T S O L S A N A N L V

CTGCTTTTGAGCAGAGCCTGGTGAATATGACATCCCGCTGCGACACC TGCGAGAGACGGCCGAGGAGAAGGACACTGAGCTGCTGGATTGCGAGAAAC
GACGAAAACCTGCTCTGGGACCACTTATACTGTAGGGCGGACGCTGTGGACCGTCTCTGCCGGCTCTCTTCTCTGTGACTCGACGACCTAAACGCTCTTTG

4100

insert pLM1

ORF pLM1

A A F E Q S L V N M T S R L R H L A E T A E E K D T E L L D L R E T

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fig 53 pLM6 (1 > 4947) Site and Sequence

Page 3

AGAAGGAGGTATCGGAGCTGCGCTCTGAGCTATGGGAGAAGGAAATGAAGCTTACAGACATCCGCTTGGAGGCCCTCAACTCTGCCCAACCAACTGGATCA
TCTTCTCCATAGCCTCGACGCGAGACTCGATACCCTCTCCCTTACTTCGAATGCTGTAGGCGAACC TCCGGGAGTTGAGACGGGTGCTTGACCTAGT

U3 stuk

ORF

K K E V S E L R S E L V E K E M K L T D I R L E A L N S A H Q L D Q

GCTTCGGGAGACCATGCACAACATGCAGTTGGAGGTGGACCTGCTGAAAGCAGAGAATGACCGACTGAAGGTAGCCCCAGGCCCTCATCAGGCTCCACT
CGAAGCCCTCTGGTACGTGTTGTACGTCAACCTCCACCTGGACGACTTTCGTCTCTTACTGGCTGACTTCCATCGGGGTCGGGGGAGTAGTCCGAGGTGA

U3 stuk

ORF

L R E T M H N M Q L E V D L L K A E N D R L K V A P G P S S G S T

CCAGGGCAGGTCCTGGATCATCTGCATTATCTTCCCCACGCCGCTCCCTAGGCCCTGGCACTCACCCATTCTTCGGCCCCAGTCTTGCAGACACAGACC
GGTCCCGTCCAGGGACCTAGTAGACGTAATAGAAGGGGTGCGGCGAGGGATCCGGACCGTGAGTGGGTAAGGAAGCCGGGGTCAGAACGCTCTGTGCTCGG

U3 stuk

ORF

P G Q V P G S S A L S S P R R S L G L A L T H S F G P S L A D T D

TGTCACCCATGGATGGCATCAGTACTTGTGGTCCAAAGGAGGAAGTGACCCCTCCGGGTGGTGGTGAGGATGCCCCCGCAGCACATCATCAAGGGGACTT
ACAGTGGGTACCTACCGTAGTCATGAACACCAGGTTTCTCTTCACTGGGAGGCCACCACCACTCTACGGGGGCGTCTGTAGTAGTTTCCCCTGAA

U3 stuk

ORF

L S P M D G I S T C G P K E E V T L R V V V R M P P Q H I I K G D L

GAAGCAGCAGGAATTCTTCTGGGCTGTAGCAAGGTCAGTGGAAAAGTTGACTGGAAGATGCTGGATGAAGCTGTTTCCAAGTGTTCAGGACTATAT
CTTCGTCTCCTTAAGAAGSACCCGACATCGTTCCAGTCACCTTTTCAACTGACCTTCTACGACCTACTTCGACAAAAGGTTCAAGTTCTTGATATAA

U3 stuk

ORF

K Q Q E F F L G C S K V S G K V D W K M L D E A V F Q V F K D Y I

TCTAAATGGACCCAGCCTCTACCTGGGACTAAGCACTGAGTCCATCCATGGCTACAGCATCAGCCACGTGAAACGAGTGTGGATGCAGAGCCCCCGS
AGATTTTACCTGGGTCGGAGATGGGACCTGATTCGTGACTCAGGTAGGTACCGATGTCTAGTTCGGTGCACTTTGCTCACAACCTACGCTCTCGGGGGG

U3 stuk

ORF

S K M D P A S T L G L S T E S I H G Y S I S H V K R V L D A E P P

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fig 53 pLM8 (1 > 4947) Site and Sequence

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AGATGCCCTCCTTGCCGTCGAGGTGTCAATAACATATCAGTCTCCCTCAAAGGTCTGAAGGAGAAATGCGTCGACAGCCTGGGTTCGAGACGCTGATCCC
TCTACGGAGGAACGGCAGCTCCACAGTTATTGTATAGTCAGAGGGAGTTTCCAGACTTCTCTTTACGCAGCTGTCGGACCACAAGCTCTGCGACTAGGS 3000

U3 stuk

ORF

E M P P C R R G V N N I S V S L K G L K E K C V D S L V F E T L I P

CAAGCCGATGATGCAGCACTACATAAGCCTCTGTGAAGCACCAGGCGCTCGTCTCTCGGGCCCCAGCGGCACGGGCAAGACCTACC TGACCAATCGG
GTTCCGGCTACTACGTCGTGATGATTTCGGAGGACGACTTCGTGGCCGCGGAGCAGGAGAGCCCGGGGTCGCCGTGCCCGTTC TGGATGGACTGGTTAGCG 3100

U3 stuk

ORF

K P M M O H Y I S L L L K H R R L V L S G P S G T G K T Y L T N R

TTGGCCGAGTACCTGGTGGAGCGCTCTGGCCGTGAGGTACAGAGGGCATCGTCAGCACCTTCAACATGCACCAGCAGTCTTGCAAGGATCTGCAACTGT
AACCAGGCTCATGGACCACCTCGCGAGACCGGCAC TCCAGTGTCTCCCGTAGCAGTCGTGGAAGTTGTACGTGGTCGTCAGAACGTTCTAGACGTTGACA 3200

U3 stuk

ORF

L A E Y L V E R S G R E V T E G I V S T F N M H O O S C K D L O L

ATCTTTCCAACCTAGCCAACAGATAGACCGGGAAACAGGAATTGGGGATGTGCCCTGGTGATTCTATTGGATGACCTGAGTGAAGCAGGCTCCATCAG
TAGAAAGGTTGGATCGGTTGGTCTATCTGGCCCTTTGTCTTAACCCCTACACGGGGACCCTAAGATAACCTACTGGACTCACTTCGTCCGAGGTTAGTC 3300

U3 stuk

ORF

Y L S N L A N O I D R E T G I G D V P L V I L L D D L S E A G S I S

TGAGTTGGTCAATGGGGCCCTCACCTGCAAGTATCATAAATGTCCCTATATTATAGGTACCACCAATCAGCCTGTAAAAATGACACCAACCATGGCTTG
ACTCAACCAGTTACCCCGGAGTGGACGTTTCATAGTATTACAGGGATATAATATCCATGGTGGTTAGTCGGACATTTTAC TGTGGGTGGTACCGAAC 3400

U3 stuk

ORF

E L V N G A L T C K Y H K C P Y I I G T T N O P V K M T P N H G L

CACCTTGAGCTTCAGGATGTTGACCTTCTCCAACAACGTGGAGCCAGCCAATGGCTTCTTGGTTCGTTACCTGAGGAGGAAGCTGGTAGAGTCAGACAGCG
GTGAACCTCGAAGTCTACAAC TGAAGAGGTTGTTGCACCTCGGTTCGGTACCGAAGGACCAAGCAATGGACTCCTCCTTCGACCATCTCAGTCTGTCCG 3500

U3 stuk

ORF

H L S F R M L T F S N N V E P A N G F L V R Y L R R K L V E S D S

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fig 53 pLM8 (1 > 4947) Site and Sequence

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ACATCAATGCCAACGAAGAGAGCTGCTCGGGTGCCTGACTGGGTACCCAAGCTGTGGTATCATCTCCACACCTTCCTTGAGAAGCACAGCACCTCAGA
TGTAGTTACGGTGTTCCTTCGACGAAGCCACGAGCTGACCCATGGGTTCGACACCATAGTAGAGGTGTGGAAGGAACCTTCGTGTCGTGAGTC 360

U3 stuk

ORF

D I N A N K E E L L R V L D V V P K L V Y H L H T F L E K H S T S D

CTTCCTCATCGGCCCTTGCTTCTTCTGTCGTGTCCTTGGCATTGAGGACTTCGGACCTGGTTCATTGACCTGTGGAACAACCTATCATTCCCTAT
GAAGGAGTAGCCGGGAACGAAGAAAGACAGCACAGGGTAACCGTAACCTCTGAAGGCTTGGACCAAGTAACGGACACCTTGTGAGATAGTAAGGGATA 370

U3 stuk

ORF

F L I G P C F F L S C P I G I E D F R T V F I D L V N N S I I P Y

CTACAGGAAGGAGCCAAGGATGGGATAAAGGTCCATGGACAGAAAGCTGCTTGGGAGGACCCAGTGGAAATGGGTCCGGGACACACTTCCTGGCCATCAG
GATGTCTTCCTCGGTTCCTACCTATTTCAGGTACCTGTCTTCGACGAACCTCTGGGTACCTTACCCAGGCCCTGTGTGAAGGGACCGGTAGTC 380

U3 stuk

ORF

L O E G A K D G I K V H G O K A A W E D P V E V V R D T L P W P S

CCCAACAAGACCAATCAAGCTGTACCACTGCCCCACCCACCGTGGGCCCTCACAGCATTGCCTCACCTCCCGAGGATAGGACAGTCAAAGACAGCAC
GGGTTGTTCTGGTTAGTTTCGACATGGTGGACGGGGTGGGTGGCACCCGGGAGTGTCTAACGGAGTGGAGGGCTCCATCTGTCAGTTTCTGTCTGTS 390

U3 stuk

ORF

A Q Q D O S K L Y H L P P P T V G P H S I A S P P E D R T V K D S T

CCCAAGTCTCTGGACTCAGATCCTCTGATGGCCATGCTGTGAACTTCAAGAAGCTGCCAACTACATTGAGTCTCCAGATCGAGAAACCATCTGGAC
GGGTTCAAGAGACCTGAGTCTAGGAGACTACCGGTACGACGACTTTGAAGTCTTCGACGGTTGATGTAACCTCAGAGGTCTAGCTCTTTGGTAGGACCTG 400

U3 stuk

ORF

P S S L D S D P L M A M L L K L O E A A N Y I E S P D R E T I L D

CCCAACCTTCAGGCAACACTTTAAGGGTTCGGCAATCACTGTACCCCCGGACAGCAGAACGCTGGCATCAGCTATCTTAGCTCTCTCTCCCTCTCTCC
GGGTTGGAAGTCGTTGTGAAATCCCAAGCCGTTAGTGACAGTGGGGGCTGTCTGCTTTCGACCGTAGTCGATAGAATCGAAGGAGAGGGAGAGG 410

U3 stuk

ORF

P N L O A T L G F G N H C H P R T A E R V H O L S L L L S P L

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fig 53 pLM6 (1 > 4947) Site and Sequence

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TCTTTCAGAGCACTGGCTCTCCAGCCCCAGGAGGAGAACAGGAGGGAGGAGGAGATGAAAGAGGAGGGACAGGTTCTTGGTGCTGTACCTTTGAGAACT~
AGAAAGTCTCGTGACCGAGAGGTCGGGGTCTCTCTTGTCTCCCTCTCTCTACTTCTCTCCCTGTCCAAGAACCACGACATGGAAACTCTTGAA

U3 stuk

L F Q S T G S P A P G G E Q E G G G D E R G G T G S V C C T F E N F

CCTAGGAAGGAATGGTGGGGTGGCGTTTGGGAAC TTGTGCCCCCTAAACACATT TAC TGGCCTCCTCTAGAGCGGCCGCCACC GCGGTGGAGCTCCAATT
GGATCCTCTCTTACCACCCACCGCAAAACCCTTGAACACGGGGGATTGTGTAATGACCGGAGGAGATCTCGCGCGGTGGCGCCACCTCGAGGTTAA

U3 stuk

L G R N G G V A F G N L C P L N T F T G L L . S G R H R G G A P I

CGCCCTATAGTGAGTCGATTACGCGCGCTCACTGGCCGTCGTTTACAACGTCGTGACTGGGAAAACCTGGCGTTACCCAACCTTAATCGCCTTGCAGC
GCGGGATATCACTCAGCATAATGCGCGCGAGTGACCGCGAGCAAAATGTTGCAGCACTGACCCCTTTGGGACCGCAATGGGTTGAATTAGCGGAACGTCG 440
R P I V S R I T R A H V P S F Y N V V T G K T L A L P N L I A L Q

ACATCCCCCTTCGCGCAGCTGGCGTAATAGCGAAGAGGCCCGCACCGATCGCCCTTCCCAACAGTTGCGCAGCCTGAATGGCGAATGGGACGCGCCCTGT
TGTAGGGGGAAGCGGTCGACCGCATTATCGCTTCFCGGGGCTGGCTAGCGGGAAGGGTGTGTAACCGCTCGGACTTACCGTTACCTGCGCGGGACA
H I P L S P A G V I A K R P A P I A L P N S C A A . M A N G T R P V

AGCGGCGCATTAAAGCGCGGCGGGTGTGGTGTACGCGAGCGTGACCGCTACACTTGCCAGCGCCCTAGCGCCCGCTCCTTTTCGCTTTCTCCCTTCC-
TCGCCCGTAATTCGCGCCGCCACACCACCAATGCGCGTCGCACTGGCGATGTGAACGGTTCGCGGGATCGCGGGCGAGGAAAGCGAAAGAAGGGAAGGA
A A H . A R R V W W L R A A . P L H L P A P . R P L L S L S S L F

TTCTCGCCACGTTGCGCGGCTTCCCGTCAAGCTCTAAATCGGGGGCTCCCTTTAGGGTTCGATTTAGTGCTTTACGGCACCTCGACCCCAAAAACT
AAGAGCGGTGCAAGCGGCCGAAAGGGCAGTTCGAGATTAGCCCCGAGGGAAATCCCAAGGCTAAATCACGAAATGCCGTGGAGCTGGGGTTTTTGA
F S P R S P A F P V K L . I G G S L . G S D L V L Y G T S T P K N

TGATTAGGGTGATGGTTACAGTAGTGGCCATCGCCCTGATAGACGGTTTTTCGCCCTTTGACGTTGGAGTCCACGTTCTTTAATAGTGGACTCTTGTTC
 ACTAATCCCACTACCAAGTGATCACCCTGGTAGCGGGACTATCTGCCAAAAAGCGGGAAACTGCAACCTCAGGTGCAAGAAATATACCTGAGAACAAC
 L I R V M V H V V G H R P D R R F F A L . R V S P R S L I V D S C S

CAAAC TGGAAACAACACTCAACCTATCTCGGTCTATTCTTTTGATTATAAGGGATTTTGCCGATTTCGGCCTATTGGTTAAAAAATGAGCTGATTAACT
GTTTGACCTTGTGTGAGTTGGGATAGACCCAGATAAGAAAACTAAATATTCCCTAAACGGCTAAAGCCGGATAACCAATTTTTACTCGACTAAATTC
K L E Q H S T L S R S I L L I Y K G F C R F R P I G . K M S . F H

AAAAAATTAAACGGAATTTTAAACAAAATTTAACGCTTACAATTTAG
 TTTTAAATTTGCGCTTAAATTTGTTTATAATTGCGAATGTTAAATC 4947
 K N L T R I L T K Y . R L Q F R

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fig 54 pLM1 (1 > 8285) Site and Sequence

Enzymes : 72 of 148 enzymes (Filtered)

Settings : Circular, Certain Sites Only, Standard Genetic Code

Page 1

GTGGCACATTTTCGGGAAATGTGCGGGAACCCCTATTGTATTATTTCTAAATACATTCAAATATGTATCCGCTCATGAGACAAATACCCCTGATAAAT 100
CAACGTGAAAAGCCCTTTACACGCGCCTTGGGGATAACAAATAAAAGATTATGTAAGTTTATACATAGGCGAGTACTCTGTTATTGGGACTATTTA
G G T F R G N V R G T P I C L F F . I H S N M Y P L M R O . P . . M
GCTTCAATAATTTGAAAAGGAAGATGAGTATTCAACATTTCCGTGTCGCCCTTATTCCTTTTTCGGGCATTTTGCCCTTCCGTGTTTGTCTCAC 200
CGAAGTTATTATAACTTTTCTTCTCACTACTAAGTTGTAAGGCACAGCGGGAATTAAGGAAAAAACGCCGTAAACGGAAGGACAAAAACGAGTG
L Q . Y . K R K S M S I Q H F R V A L I P F F A A F C L P V F A H
CCAGAAACGCTGGTGAAGTAAAGATGCTGAAGTACAGTTGGGTGCACGAGTGGGTACATCGAACTGGATCTCAACAGCGGTAAAGATCCTTGAGAGTT 300
GGTCTTTGGCACCCTTTCAATTTTCTACGACTTCTAGTCAACCCACGTGCTCACCCTAATGTAAGTTGACCTAGAGTTGTCGCCATTCTAGGAACCTCTCAA
P E T L V K V K D A E D O L G A R V G Y I E L D L N S G K I L E S
TTGCCCCGAAGAAGCTTTTCAATGATGAGCACTTTTAAAGTTCTGCTATGTGGCGCGGTATTATCCCGTATTGACGCCGGCAAGGCAACTCGGTG 400
AAGCGGGGCTTCTGCAAAAGGTTACTACTCGTGAAATTTCAAGACGATACACCGGCCATAATAGGCGATAACTGCGGCCGTTCTGTTGAGCCAGC
F R P E E R F P M H S T F K V L L C G A V L S R I D A G O E O L G R
CCGCATACACTATTCTCAGAATGACTTGGTTGAGTACTCACCAGTCACAGAAAAGCATCTTACGGATGGCATGACAGTAAGAGAATTATGCAGTGTGCC 500
GGCGTATGTGATAAGAGTCTTACTGAACCACTCATGAGTGGTCAGTGCTTTTCTAGTAAGTCCCTACCGTACTGTCTTCTTAATACGTCACGACGG
R I H Y S Q N D L V E Y S P V T E K H L T D G M T V R E L C S A A
ATAACCATGAGTGATAACACTGCGGCCAATCTACTTCTGACAACGATCGGAGGACCGAAGGAGCTAACCGCTTTTTCACAACTGGGGGATCATGTAA 600
TATTGGTACTCACTATTGTGAGCGCGGTTGAATGAAGACTTGTGCTAGCTCCTGGCTTCTCGATTGGCGAAAAACGTGTGTACCCCTAGTACATT
I T M S D N T A A N L L L T T I G G P K E L T A F L H N M G D H V
CTGCGCTTGATCGTTGGGAACCGGAGCTGAATGAAGCATACCAACGACGAGCGTGACACCAGATGCCGTAGCAATGGCAACAACGTTGCGCAAACT 700
GAGCGGAATAGCAACCTTGGCTCGACTTACCTCGGTATGGTTTGTGCTGCGACTGTGGTGTACGGACATCGTTACCGTTGTGCAACGCGTTTGA
T R L D R V E P E L N E A I P N D E R O T T M P V A M A T T L R K L
ATTAACGCGCACTACTTACTCTAGCTTCCCGCAACAATTAATAGACTGGATGGAGGCGGATAAAGTTGCAGGACCACCTTCTGCGCTCGGCCCTTCCG 800
TAATTGACCGCTTGATGAATGAGATCGAAGGCGGTTGTAATTATCTGACCTACCTCCGCTATTTCACGTCCTGGTGAAGACGCGAGCCGGGAAGGC
L T G E L L T L A S R Q O L I D V H E A D K V A G P L L R S A L P
GCTGGCTGGTTTATTGCTGATAAATCTGGAGCCGTTGAGCGTGGGTCTCGCGGTATCATTCGACACTGGGGCCAGATGGTAAGCCCTCCCGTATCGTAG 900
CGACCGACCAATAACGACTATTTAACCTCGGCCACTCGCACCCAGAGCGCCATAGTAACCTCGTGACCCCGGTCTACCAATCGGGAGGCGATAGCATC
A G V F I A D K S G A G E R G S R G I I A A L G P D G K P S R I V
TTATCTACACGAGGGGAGTCAGGCAACTATGGATGAACGAAATAGACAGATCGCTGAGATAAGTCCCTACCTGATTAAAGCATTGGTAACGTGACACCA 1000
AATAGATGCTGCCCTCAGTCCGTTGATACCTACTTGTCTTATCTGCTAGCGACTCTATCAGCGAGTGACTAATTCGTAACCATTGACAGTCTGGT
V I Y T T G S Q A T M D E R N R O I A E I G A S L I K H V . L S D Q
AGTTTACTCATATAFACCTTTAGATTGATTTAAACTTCAATTTTAAATTTAAAGGATCTAGGTAAGATCCTTTTGTATAATCTCATGACCAAAATCCCT 1100
TCAATGAGTATATAGAACTAATAAATTTGAAGTAAATAAATTTTCTTAGATCCCTTCTAGGAAAACTATTAGAGTACTGGTTTATAGGGA
V Y S Y I L . I D L K L H F . F K R I . V K I L F D N L M T K I P
TAAGTGAGTTTTCGTTCCACTGACGCTGAGACCCGCTAGAAAAAGATCAAGGATCTCTTGAATCCTTTTCTGCGGTAATCTGCTGCTTGCAAA 1200
ATTGCACTCAAAAGCAAGGTGACTGCACTGCGGCTCTTTTCTAGTTTCTAGAAGAACTCTAGGAAAAAAGACGCGCATTAGACGACGAACGTTT
. R E F S F H . A S D P V E K I K G S S . D P F F L R V I C C L O
CAAAAAACCAACGCTACGACGCTGGTTGTTTGGCGGATCAAGAGCTACCAACTCTTTTTCGAAGGTAACGGCTTACGACGAGCGCAGATACCAAA 1300
GTTTTTTTGGTGGGATGGTGGCCACCAACACCGGCTAGTTCTCGATGGTGTAGAAAAAAGTCTTCCATTGACCGAAGTCGCTCGCGCTCTATGGTTT
T K K P P L P A V V C L P D O E L P T L F P K V T G F S R A Q I P N
TACTGCTCTCTAGTGTAGCGGTAGTTAGGCCACCACTTCAAGAACTCTGTGACACCGCTCTATACCTCGCTCTGCTAATCTGTTACCAAGTGGCTGCT 1400
ATGACAGGAAGATCAGATCGGATCAATCCGGTGGTGAAGTTCTTGAGACATCGTGGCGGATTTATGGAGCGAGACGATTAGGACAAATGGTACCGACGA
T V L L V . P . L G H H F K N S V A P P T Y L A L L I L L P V A A

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GCAGTGGCGATAAGTCGTGCTTACCGGTTGGACTCAAGACGATAGTTACCGGATAAGGCGCAGCGGTCGGGCTGAACGGGGGGTTCGTGCACACAGC
CGGTCAACCGCTATTCAGCAGAGAATGGCCCAACCTGAGTTCGTATCAATGGCTATTTCCGCGTCGCCAGCCGACTTGCCCCCAAGCACGTGTGTGC 1500
A S G D K S C L T G L D S R R . L P D K A Q R S G . T G G S C T O

CCAGCTTGGAGCGAACGACCTACACCGAATGAGATACCTACAGCGTGAGCTATGAGAAAGCGCCAGCTTCCGAAGGGAGAAAGGCGGACAGGTATCC
GGTCGAACCTCGCTTGGATGTGGCTTGACTCTATGGATGTCGCACTCGATACCTTTCCGGTGCGAAGGGCTTCCCTCTTTCCGCTGTCCATAGG 1600
P S L E R T T Y T E L R Y L O R E L . E S A T L P E G R K A D R Y P

GGTAAGCGCGAGGGTCGGAACAGGAGAGCGACGAGGGAGCTTCCAGGGGGAACGCCGTGGTATCTTTATAGTCTGTGGGTTTCGCCACCTCTGACTT
CCATTCGCGCTCCCGAGCTTGTCTCTCGCGTGTCTCTCGAAGGTCCCTTTGCGGACCATAGAAATATCAGGACAGCCAAAGCGGTGGAGACTGAA 1700
V S G R V G T G E R T R E L P G G N A V Y L Y S P V G F R H L . L

GAGCGTCGATTTTGTGATGCTCGTCAGGGGGCGGAGCCTATGGAACACGCCAGCAACCGGGCTTTTACGGTTCCTGGCTTTTGTGGCTTTTG
CTGCGAGCTAAACACACTACGAGCAGTCCCCCGCTCGGATACCTTTTTCGGTTCGTTCGCGCGGAAATGCCAAGGACCGGAAACGACGGAAC 1800
E R R F L . C S S G G R S L V K N A S N A A F L R F L A F C V P F

CTCACATGTTCTTCTCGCTTATCCCTGATTCTGTGGATAACCGTATTACCGCTTTGAGTGAGCTGATACCGCTCGCCGAGCGAAGCAGCGAGCG
GAGTGACAAAGAGCGCAATAGGGACTAAGACACCTATTGGCATAATGGCGGAACTCACTCGACTATGGGAGCGGCTCGGCTTGTGGCTCGC 1900
A H M F F P A L S P D S V D N R I T A F E . A D T A R R S R T T E R

CAUCGAGTCAGTGAGCGAGGAAGCGGAAGAGCGCCCAATACGCAACCGCTCTCCCGCGTGGGCGATTTCATTAATGACGCTGGCAGCAGGTTT
GTCGTCAGTCACTCGCTCTCTCGCTTCTCGCGGTTCGCGGAGAGGGGCGCAACCGCTTAAGTAATTACGTCGACCGTGTGTCCAAA 2000
S E S V S E E A E E R P I R K P P L P A R V P I H . C S V H D R F

CCGAC TGGAAAGCGGCGAGTGAGCGCAACGAATTAATGAGTTAGCTCACTATTAGGACCCAGGCTTTACACTTTATGCTTCGGCTCGTATGT
GGGCTGACCTTTCCCGCTCACTCGGTTGGCTTAATTACACTCAATCGAGTGAGTAATCCGTGGGTCGGAATGTGAATACGAAGGCCGAGCATACA 2100
P D V K A G S E R N A I N V S . L T H . A P Q A L H F H L P A R M

TGTGTGAATTGTGAGCGGAATAACAATTTACACAGGAACAGCTATGACCATGATTACGCCAAGCGCGCAATTAACCTCACTAAAGGAACAAAGCT
ACACACCTTAACACTCGCTTATGTTAAAGTGTCTCTTTCGATACCTGCTACTAATGCGGTTTCGCGCTTAATTGGGAGTGATTCCCTTGTTCGA 2200
L C G I V S G . O F H T G N S Y D H D Y A K R A I N P H . R E Q K L

GGGTACCGGGCCCCCTCGAGGTCGACGGTATCGATAAGCTTGAATATCGAATCTCGAGCCCTGCTCTTCAGCCAGATGCTGGACCCAGAGTCCAG
CCATATGGCCCGGGGGGAGCTCCAGCTGCCATAGCTATTGAACTATAGCTTAAGGAGCTCGGGGACGAGAAGTCGGTCTACGACCTGGGTCTCAGGGTC 2300

insert pLM1

ORF pLM1

G T G P P L E V D G I D K L D I E F L Q P L L F S Q M L D P E S O

AGAAAGAGGACAGTGCAGAAATGCTCTGGATCTCCGGCAGAACC TGGAAAGAGCAATGTCAGCTCGGAGGGTCCAGGTGACTCACAGCTCCCTGGAGA
TCITTCCTCTGTCAGCTCTTACAGGACCTAGAGGCGCTTGGACCTTCTCTGGTACAGGTGCGGACGCTCCAGGGTCCACTGAGTGTGAGGGACCTCT 2400

insert pLM1

ORF pLM1

R K R T V O N V L D L R Q N L E E T M S S L R G S Q V T H S S L E

TGACCTGCTACGACAGCGATGATGCCAACCCACGAGCGTGTCAGCTCTCCAACCGCTCGTCCCTCTGTATGGCGCTATGGCCAGTCCAGTCCGCG
ACTGGACGATGCTGCTGCTACTACGGTTGGGTGCGTCGCACAGGTGCGAGAGGTTGGCGAGCAGGGGAGACAGTACCGGATACCGGTACAGTACGGCGC 2500

insert pLM1

ORF pLM1

H T C Y D S D D A N P R S V S S L S N R S S P L S V R Y G C S S P R

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fig 54 pLM1 (1 > 8285) Site and Sequence

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GC TGCAGGCTGGTGACGCGCCCTCTGTGGG TGGGAGC TGCCTGCTGGAGGGGACGCCGCTTGGTACATGCACGGGGAACGGGCCACTATCCACACC
CGACGTCCGACCACTGCGCGGGAGACACCCACCTCGACGGCGAGCCTCCCTGCGGGGGACCAATGTACGTGCCCTTGGCCGGGTGATGAGGGTGTGG 2800

— insert pLM1 —
— ORF pLM1 —
L Q A G D A P S V G G S C R S E G T P A V Y M H G E R A H Y S H T

ATGCCCATGCCAGCCCCAGCAAGCTCAGCCATATCTCCCGCTGGAGCTGGTCGAATCCCTGGACTCGGATGAGGTGGACCTCAAGTCCGGCTACATGA
TACGGGTACGGTCGGGGTCGTTTCGAGTCGGTATAGAGGGCGGACCTCGACCACTTAGGGACCTGAGCCTACTCCACCTGGAGTTCAGGCCGATGTACT 2700

— insert pLM1 —
— ORF pLM1 —
M P M R S P S K L S H I S R L E L V E S L D S D E V D L K S G Y M

GCGACAGTGACCTCATGGCAAGACCATGACGGAGGATGATGACATCACTACCGGCTGGGATGAAAGCAGCTCCATCAGTAGTGGACTCAGCGATGCCCTC
CGCTGTCACTGGAGTACCGGTTCTGGTACTGCCCTCTACTACTGTAGTGATGGCCGACCTACTTTGTCGAGGTAGTCATCACCTGAGTCGCTACGGAG 2800

— insert pLM1 —
— ORF pLM1 —
S D S D L M G K T M T E D D D I T T G V D E S S S I S S G L S D A S

AGACAATCTCAGTTCAGAAGAATTCAATGCCAGCTCCTCACTCAACTCCCTCCCAAGTACTCCCACTGCTTCTCGCAGGAACCAACAATAGTGTACGC
TCTGTTAGAGTCAAGTCTTCTTAAGTACGGTCGAGGAGTGAGTTGAGGGAGGGTTCATGAGGGTGACGAAGAGCGTCTTGAGTTGTTATCACGATGCG 2900

— insert pLM1 —
— ORF pLM1 —
D N L S S E E F N A S S S L N S L P S T P T A S R R N S T I V L R

ACAGACTCAGAGAAGCGCTCACTGGCAGAAAGTGGGCTGAGCTGGTTAGTGAAATCAGAGGAGAAAGCCCCATAAAAACTGGAGTACGACAGTGGTAGCC
TGCTGAGTCTCTTCGCGAGTGACCGCTCTTCAACCGGACTCGACCAAACTACTAGTCTCTCTTTGCGGGGATTTTGTGACCTCATGCTGTCACCATCGG 3000

— insert pLM1 —
— ORF pLM1 —
T D S E K R S L A E S G L S V F S E S E E K A P K K L E Y D S G S

TGAAGATGGAACCTGGGACTTCTAAGTGGCGGAGGGAGCGGCC TGAGAGCTGTGATGATTCATCCAAGGGTGGAGAATGAAAAAGCCCATCAGCCTGGG
ACTTCTACCTTGGACCTGAAGATTCACCGCTCCCTCGCCGGACTCTCGACACTACTAAGTAGGTTCACCTCTTGAC TTTTTCGGGTAGTCGGACCC 3100

— insert pLM1 —
— ORF pLM1 —
L K M E P G T S K V R R E R P E S C D D S S K G G E L K K P I S L G

CCACCTGGTTCCCTGAAGAAGGGCAAGACCCACCTGTGGCTGTAACCTCCCCATCACTCACACGCCAGAGTGGCTCAAAGTCGAGGCAACCT
GGTGGGACCAAGGACTTCTTCCCGTTCGGGGTGGACACCGACATTGAAGGGGGTAGTGAGTGTGTCGGGTCTCACGGGAGTTTCAGCGTCCGTTTGA 3200

— insert pLM1 —
— ORF pLM1 —
H P G S L K K G K T P P V A V T S P I T H T A Q S A L K V A G K P

GAGGGCAAAGCTACAGACAAGGTAAGCTTGCAGTGAAGAATCTGGGCTCAACGCTCTCTCTGTGATGCTGGTGGGACCGCTGAGTGATGCTAAGA
CTCCCGTTTCGATGCTGTTCCTTCGAACTCAAGCTCACTTCTTATGACCCGAGGTTCGAGGAGGAGACTACGACAGCCCTGGCGGACTCACTACGATCT 3300

— insert pLM1 —
— ORF pLM1 —
E G K A T D K G K L A V K N T G L O R S S S D A G R D R L S D A K

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AGCCCCCTCGGGCATTGCTCGCCCCCTCCACTTCGGGATCCCTCGGCTACAAGAAGCCCTCCCTGCCACAGGCACAGCCACTGTCATGCAAACTGGTGG 3400
TCGGGGGAGCCCGTAACGAGCGGGGAGGTGAAGCCCTAGGAAGCCGATGTTCTCGGAGGAGGACGGTGTCCGTGTCGGTGACAGTACGTTTGACCACT
-----insert pLM1-----
-----ORF pLM1-----
K P P S G I A R P S T S G S F G Y K K P P P A T G T A T V M Q T G G
TTCAGCCACTCTCAGCAAGATCCAGAAGTCTCAGGCATCCCTGTCAAGCCAGTAAATGGGCGCAAGACTAGCTTAGAGTGTTCACAGCGCAGAGCCA 3500
AAGTCGGTGAGAGTCGTTCTAGGTCTTCAGGAGTCCGTAGGGACAGTTCGGTCATTTACCCGCGTTCGTGCAATCTACAAAGGTTGTCGGCTCCTGGT
-----insert pLM1-----
-----ORF pLM1-----
S A T L S K I O K S S G I P V K P V N G R K T S L D V S N S A E P
GGATTCTCGGCTCTGGAGCCCGTCTCAACATCCAGTACCGCAGCCCTGCCCCGGCCAGCCAAAGTCAAGTTCTATGAGCGTGACCGCGGGGGGGTGGAC 3600
CCTAAGGACCGAGGACCTCGGGCAAGATTGTAGGTCAATGGCTCGGACGGGGCCGGTTCGGTTCAGTCAAGATACTCGCACTGGCCCGCCGCCCACTGT
-----insert pLM1-----
-----ORF pLM1-----
G F L A P G A R S N I Q Y R S L P R P A K S S S H S V T G G R G G
CTCGCCCTGTGAGCAGCAGCATTGACCCAGTCTCTCAGCACCAAGCAGGAGGGCCCTACGCCCTTCAGACTGAAGGAGCCCTACCAAGGTAGCCAGTGG 3700
GAGCGGGACACTCGTCTGTAACCTGGGGTCAGAGGAGTCGTGGTTCGTCCTCCGGAATCGGGAAGGTCGACTTCCTCGGATGGTTCATCGGTCACC
-----insert pLM1-----
-----ORF pLM1-----
P R P V S S S I D P S L L S T K Q G G L T P S R L K E P T K V A S G
GCGGACCACTCCAGCCCTGTCAATCAGACAGATCGGGAAAAGGAGAAGGCCAAAGCCAAAGCAGTGGCTTGGACTCAGACAACATCTCCTTGAAGAGT 3800
CGCCCTGGTGAGGTCGGGGACAGTTAGTCTGTCTAGCCCTTTCTCTTCCGGTTTCGGTTCGCTACCGGAACCTGAGTCTGTTGAGAGGAACCTCTCA
-----insert pLM1-----
-----ORF pLM1-----
P T T P A P V N Q T O R E K E K A K A V A L D S D N I S L K S
ATTGGCTCCCCAGAGGACTTCCCAAGAACAAGCAAGCCACCCACAGCCACCAAGCTGGCAGAGCTGCCAACCCCTCTCAGGGCCACAGCGAAGA 3900
TAACCGAGGGGCTCTCATGAGGGTCTCTGGTTCGTTCCGTTGGGTCGCGTGGTTCGACCTCTCAGCGGTGGTGGGGAGAGTCCCGGTGTCGCTTCT
-----insert pLM1-----
-----ORF pLM1-----
I G S P E S T P K N Q A S H P T A T K L A E L P P T P L R A T A K
GTTTGTCAAAACCCCTCACTAGCCAATCTTGACAAGGTCAACTCCAAAGCTCTGGATCTACCATCATCCAGTGATACCCCATGCTTCAAGGTCCC 4000
CJAACAGTTTGGTGGGAGTGATCGGTAGAACGTTCAGTTGAGGTTGTGAGACCTAGATGGTAGTAGGTCACTATGGTGGGTACGAAGTTTCCAGGG
-----insert pLM1-----
-----ORF pLM1-----
S F V K P P S L A N L O K V N S N S L D L P S S S D T T H A S K V P
AGATCTGATGCTACAAGCTCAGCATCTGGGGCCCTCTCCCTTCCTGCTTACCCCCAGTCCGGCACCCATCTCAATATTAACCTAGCGAGCTTCTCC 4100
CTAGACGTACGATGTTGAGTCGTAGACCCCGGAGAGGGAAGGACGAAGTGGGGTTCAGGCCGTGGTAGGAGTTATAATTGAGTCGGTCAAGAGG
-----insert pLM1-----
-----ORF pLM1-----
U L H A T S S A S G G P L P S C F T P S P A P I L N I N S A S F S

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fig 54 pLM1 (1 > 8285) Site and Sequence

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CAGGGCCCTGGAGCTAATGAGTGGTTCAGTGTGCCAAAAGAGACCCGCATGTACCCCAAACCTCAGGGCCTGCACAGGAGCATGGAGTCCCTCCAGATGC 4200
GTCCCGGACCTCGATTACTACCAAAGTCACACGGTTTCTCTGGGCGTACATGGGTTTGAGAGTCCGGACGTGTCTCGTACCTCAGGGAGGCTACG

— insert pLM1 —

ORF pLM1

Q G L E L M S G F S V P K E T R M Y P K L S G L H R S M E S L O M

CAATGAGCCTCCCAAGTGCCTTCCCAAGCAGTACTCCCGTCCCAACCCACCTGCTCCCAAGTGCCTCAGAGAAGAGAGACGGAAGAGCTGACTTG 4300
GTACTCGGAGGGGTCACGGAAGGGTCTCATGAGGGCAGGGGTGGGGTGGACGAGGGGACGACGAGGGTGTCTTCTCTGCTCTGCTGACTGAAC

— insert pLM1 —

ORF pLM1

P M S L P S A F P S S T P V P T P P A P P A A P T E E E T E E L T V

GAGTGAAGCCCAAGAGCTGGGCAAC TGGACAGTAATCAGCGGGATCGGAACACTCTTCCCAAGAAAGGGCTCAGGTACACAGCTTCAGTCCCAGGAGGAG 4400
CTCACCTTCGGGGTCTCGACCCGTGACCTGTCTATTAGTCGCCCTAGCCTGTGAGAAGGGTCTTCTCCGAGTCCATGGTCGAAGTCAGGGTCCCTCTC

— insert pLM1 —

ORF pLM1

S G S P R A G O L D S N O R O R N T L P K K G L R Y O L O S Q E E

ACCAAGGAGAGGGCAGATTCCCAATACCATTTGGTGGGCTGCCGTAATCCGATGACAGTACAGAGCTGCCTTCTCCCTTGCCTTCCCATGCTCTGAGTG 4500
TGGTTCCTCTCCGCTGTAAGGGTATGGTAACCAACCCGACGGACTTAGGCTACTGGTCACTCTGACGGAAGAGGGGACGTGAAGGGTACAGAGACTCAC

— insert pLM1 —

ORF pLM1

T K E R R H S H T I G G L P E S D D O S E L P S P P A L P M S L S

CAAAGGGCCAACCTTACCAACATAGTGAGTCCCACTGCGGCCACCAAGCAAGCAATCCCAAGCAATCCCAAGCAAGCAAGCAAGCAAGCAAGCAAGCAAG 4600
GTTTCCCGGTGAATGGTTGATACATCAGGGTACGCGGCTGGTGGGTTCTTAGTGGGCGAGGTGTCTGAGGGGTGGGTGCTCCGCGGAAGCTCGA

— insert pLM1 —

ORF pLM1

A K G O L T N I V S P T A A T T P R I T R S N S I P T H E A A F E L

GTACAGCGGCTCCCAATGGGAGCACCCTGCTCCGCGGAGAGACCAAGGGAATGATTCGGTCAGGATCCTTCCGAGACCCACGGACGATGTTTAC 4700
CATGTGCGCTAGGGTTTACCCCTCGTGGGACAGGACCGGCTCTCTGGGTTCCCTTACTAAGCCAGTCTAGGAAGGCTCTGGGGTGGCTGTACAAGTG

— insert pLM1 —

ORF pLM1

Y S G S O M G S T L S L A E R P K G M I R S G S F R D P T D V H

GGCTCAGTGCTGCTCCCTGGGCTCCAGTGCCTCTCCACCTACTCTCAGCTGAGGAGAGGATGCAATCTGAGCAATCCGGAAGCTTCGTAGGGAAGTGG 4800
CCGAGTCACGACAGGACCGGAGGTACGAGGAGGTGGATGAGGAGTGCAGTCTCTCTACGTTAGACTCGTTTAGGCTTCGAAGCATCCCTTGACC

— insert pLM1 —

ORF pLM1

G S V L S L A S S A S S T Y S S A E E R M O S E O I R K L R R E L

AATCATCCCAAGGAAAGTGGCCACCTTGACGTCTCAGCTTCTGCAATGCTAATCTGGTGGCTGCTTTTGGAGCAGGCTGGTGAATATGACATCCG 4900
TTAGTAGGGTCTCTTTTACCGGTGAAC TGCAGAGTCGAAAGACGGTTACGATTAGACACCGACGAAACCTGCTCGGACCACTTATACGTAGGGC

— insert pLM1 —

ORF pLM1

E S S O E K V A T L T S O L S A N A N L V A A F E O S L V N M T S R

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CCTGCGACACCTGGCAGAGACGGCCGAGGAGAAGGACACTGAGCTGCTGGATTTCGAGAGAACCATAGACTTTCTGAAGAAAAGAACCTGAGGCCAG
GGACGCTGTGGACCGTCTCTGCGGCTCCTCTCTCTGTGACTCGACGACCTAAACGCTCTTTGGTATCTGAAAGACTTCTTTTCTTGAGACTCCGGGTC 5000
-----
insert pLM1
-----
ORF pLM1
-----
L R H L A E T A E E K D T E L L O L R E T I D F L K K K N S E A Q
GCAGTCATTCAGGGAGCCCTTAATGCCTCAGAAACACACCCAAAGAACCTCGGATCAAGAGACAAAACCTCCAGATAGCATCTCAAGCCTCAACAGCA
CGTCAGTAAGTCCCTCGGAATTACGGAGTCTTTGGTGTGGGTTCTTGAAGCCTAGTCTCTGTTTGGAGGAGTCTATCGTAGAGTTCGGAGTGTCTGT 5100
-----
insert pLM1
-----
ORF pLM1
-----
A V I Q G A L N A S E T T P K E L R I K R O N S S D S I S S L N S
TCAC TAGCCATTCAGCATCGGCAGCAGCAAGGATGCTGATGCGAAAAGAAGAAAAAAGAGTTGGGTCATGAGCTTCGAAGTTCCTTCAACAAAGC
AGTGATCGGTAAGGTCGTAGCCGTGCTGCTCTCTACTACGCTTTTCTCTTTTCTCTCAACCCAGATACCTGAAGCTTCAAGGAAGTGTGTCG 5200
-----
insert pLM1
-----
ORF pLM1
-----
I T S H S S I G S S K D A D A K K K K K K S V V Y E L R S S F N K A
GTYCAGTATAAAAAAGGGGCCCAAGTCAGCTTCTCTCATCTCGGATATAGAGGAGATTGCTACACCCGACTCTTCAGCCCTCATCCCCAACTACAG
CAAGTCATATTTTCCCGGGGTCAGTCGAAGGAGTATGAGCCTATATCTCTCTTAACGATGTTGGGCTGAGAAGTCGGGGGAGTAGGGGGTTGATGTC 5300
-----
insert pLM1
-----
ORF pLM1
-----
F S I K K G P K S A S S Y S D I E E I A T P D S S A P S S P K L Q
CATGGTTCCACAGAGACTGCTTCAACCTCCATCAAGTCTCCACCTTGTCTCTCCGTGGGCACTGATGTCACCGAGGGCCCTGCTCACCAGCCCCCACA
GTACCAAGGTGCTCTGACGAAGTGGGAGGTAGTTCAGGAGTGGAAACAGGAGGCACCCGTGACTACAGTGGCTCCCGGGACGAGTGGGTCGGGGGGTGT 5400
-----
insert pLM1
-----
ORF pLM1
-----
H G S T E T A S P S I K S S T L S S V G T D V T E G P A H P A P H
CTAGGCTGTTCATGCAAAAGAGGAGGAGGAGCAGAGAAGAAGGAGGTATCGGAGCTGCGCTCTGAGCTATGGGAGAAGGAAATGAAGCTTACAGACAT
GATCCGCAAGGTACGTTTACTCTCTCTCTCGGTCTCTCTCTCTCA TAGCTCGACGCGAGACTCGATACCTCTCTCTTACTTCGAAATGCTGTGA 5500
-----
insert pLM1
-----
ORF pLM1
-----
T R L F H A N E E E E P E K K E V S E L R S E L V E K E M K L T D I
CCGC TTGGAGGCCCTCAACTCTGCCACCAACTGGATCAGCTTCTGGGAGACCATGCACAACATGCAGTTGGAGGTGGACC TGCTGAAAGCAGAGAATGAC
GGCGAACC TCCGGAGTTGAGACGGGTGGTIGACCTAGTCGAAGCCCTCTGGTACGTGTTGTACGTCAACCTCCACCTGGACGACTTTCGTCCTTACTG 5600
-----
insert pLM1
-----
ORF pLM1
-----
R L E A L N S A H Q L D O L R E T H H N M Q L E V O L L K A E N D
CGACTGAAGGTAGCCCCAGGCCCTCATCAGGCTCCACTCCAGGGCAGGTCCCTGGATCATCTGCATTATCTTCCCCACGCCGCTCCCTAGGCCGTGGAC
CTGACTTCCATCGGGGTCGGGGAGTAGTCCGAGGTGAGGTCCCTCCAGGGACCTAGTAGACGTAATAGAAGGGGTGCGGCGAGGGATCCGGACCGTG 5700
-----
insert pLM1
-----
ORF pLM1
-----
R L K V A P G P S S G S T P G O V P G S S A L S S P R R S L G L A
```

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fig 54 pLM1 (1 > 8285) Site and Sequence

Page 1

TCACCCATTCCTTCGGCCCCAGCTTCGACACACAGACCTGTCACCCATGGATGGCATCAGTACTTGTGGTCCAAAGGAGGAAGTGACCTTCGGGTGGT
AGTGGGTAAGGAAGCCGGGTCAGAACGCTGTGTCTGGACAGTGGGTACCTACCGTAGTCATGAACACCAGGTTTCTTCCTCACTGGGAGGCCACCA 5800

insert pLM1

ORF pLM1

L T H S F G P S L A D T D L S P M D G I S T C G P K E E V T L R V V
GGTGAGGATGCCCCGAGCACATCATCAAGGGGACTTGAAGCAGCAGGAATTCCTTGGGCTGTAGCAAGGTCACTGGGAAAGTTGACTGGAAGATG
CCACTCTACGGGGCGTGTGTAGTAGTTTCCCTGAACTTCGTCGTCTTAAGAAGGACCCGACATCGTTCAGTCACCTTTCAACTGACCTTCTAC 5900

insert pLM1

ORF pLM1

V R M P P Q H I I K G D L K Q Q E F F L G C S K V S G K V D V K H
CTGGATGAAGCTGTTTCCAAGTGTCAAGGACTATATTCTAAAATGGACCCAGCCTCTACCTGGGACTAAGCACTGAGTCCATCCATGGCTACAGCA
GACCTACTTCGACAAAGGTTCAAGTTCCTGATATAAAGATTTACCTGGGTCGGAGATGGGACCTGATTCTGTGACTCAGGTAGGTACCGATGTCGT 6000

insert pLM1

ORF pLM1

L D E A V F Q V F K D Y I S K M D P A S T L G L S T E S I H G Y S
TCAGCCACGTGAAACGAGTGTGGATGCAGAGCCCCGAGATGCTCTTTCGCGTCGAGGTGTAATAACATATCAGTCTCCCTCAAAGGTCTGAAGGA
AGTCTGGTGCACTTTGCTCACAACCTACGTC TCGGGGGGCTCTACGGAGGAAACGGCAGCTCCACAGTTATTGTATAGTCAGAGGGAGTTTCCAGACTTCT 6100

insert pLM1

ORF pLM1

I S H V K R V L D A E P P E M P P C R R G V N N I S V S L K G L K E
GAAATGCGTCGACAGCCTGGTGTTCGAGACGCTGATCCCCAAGCCGATGATGACGACTACATAAGCCTCTGCTGAAGCACGGGCGCTCGTCTCTCG
CTTACGCACTGTCCGACCACAAGCTCTGCGACTAGGGGTTCGGCTACTACGTCGTGATGATTTCGGAGGACGACTTCGTGGCCGCGGAGCAGGAGAGC 6200

insert pLM1

ORF pLM1

K C V D S L V F E T L I P K P M H O H Y I S L L L K H R R L V L S
GGCCCCAGCGGCACGGCAAGACCTACCTGACCAATCGCTTGGCCGAGTACCTGGTGGAGCGCTCTGGCCGTSAGGTACAGAGGGCATCGTCAGCACCT
CGUUGGTCGCCGTGCCGTTCGGATGGACTGGTTAGCGAACCGGCTCATGGACCACTCGCGAGACCGGCATCCAGTGTCTCCCGTAGCAGTCGTGGA 6300

insert pLM1

ORF pLM1

G P S G I G K T Y L T N R L A E Y L V E R S G R E V T E G I V S T
TCAACATGCACCAAGCTCTGCAAGGATCTGCAACTGTATCTTTCCAACCTAGCCAACAGATAGACCGGGAACAGGAATTGGGGATGTGCCCTGGT
AGTTGTACGTGGTGTGAGAACGTTCTAGACGTTGACATAGAAAGTTGGATCGGTTGGTCTATCTGGGCCCTTTGTCTTAACCCCTACACGGGGACCA 6400

insert pLM1

ORF pLM1

F N M H O O S C K D L O L Y L S N L A N O I D R E T G I G D V P L V
GATTCTATTGGATGACCTGAGTGAAGCAGGCTCCATCAGTGAGTTGGTCAATGGGGCCCTCACCTGCAAGTATCATAAATGTCCCTATATTATAGGTACC
CTAAGATAACCTACTGGACTCACTTCGTCCGAGGTAGTCACTCAACAGTTACCCCGGAGTGGACGTTCATAGTATTACAGGGAATAATATCCATGG 6500

insert pLM1

ORF pLM1

I L L D D L S E A G S I S E L V N G A L T C K Y H K C P Y I I G T

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[illegible]

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fig 54 pLM1 (1 > 8285) Site and Sequence

Page 1

AGGAGGGACAGGTTCTTGGTGCTGTACCTTTGAGAACCTCC TAGGAAGGAATGGTGGGGTGGCGTTTGGGAACCTTGTCCTCCCTTAAACACATTTACTGGC
TCTTCCCTGTCCAAGAACCACGACATGGAACTCTTGAAGGATCTTCTTACCACCCACCGCAAACCTTGAACACGGGGGATTGTGTAAATGACCG 7400

insert pLM1

G G T G S V C C T F E N F L G R N G G V A F G N L C P L N T F T G

CTCTCTAATGACTTTGGGAAAAGATGATTCTGGGTCTTCCCTTGACTTCTTGTTCAATTACAACTCTGGGCTTCTGGGGAGGGGTTTCAGAAAA 7500
GAGGAGATTACTGAAACCCCTTTTCTACTAAGACCCAGAAAGGGAACGAAGAACAAGTTAATGTTTGAGGACCCGAAAGACCCCTCCCAAGTCTTTT

insert pLM1

L L . . L V G K D D S G S F P . L L V S I T N S V A F V G G V Q K

CATCAAACTACTGACAGTTCCTCCGAATTCAGCTTGACTTAACCAAGGCTGAACCTTGCTCAAAAGAGCCGAATTCAGCACACTGGCGGCGTTACT 7600
GTAGTTTGTGACGTCGTAAGGGGCTTAAGTCGAACCTGAATTTGGTCCGACTGAACGAGTTTCTTCGGCTTAAGTCGTCGTGACCGCGGCAATGA

insert pLM1

T S K H C S S S P E F S L D L T R L N L L K R S R I P A H V R P L L

AGTTCTAGAGCGGCGCCACCGCGGTGGAGCTCAATTGCGCCTATAGTGAGTCGATTACGCGGCTCACTGGCGGCTGTTTACAACGTCGTGACTGG 7700
TCAAGATCTCGCGGCGGTGGCGCCACCTCGAGGTTAAGCGGGATACACTCAGCATAATGCGCGGAGTGACCGGACGAAATGTTGACGACTGACC

→

V L E R P P P R V S S N S P Y S E S Y Y A R S L A V V L O R R D V

GAAACCTGGGTTACCAACTTAATCGCTTGACGACATCCCTTTTCGCCAGCTGGCGTAATAGCAAGAGGCCCGACCGATCGCCCTTCCCAAC 7800
CTTTTGGGACCGCAATGGTTGAATTAGCGGAACGTCGTGTAGGGGGAAAGCGGTGACCGCATTATCGCTTCTCCGGGCTGGCTAGCGGGAAGGGTTG

E N P G V T Q L N R L A A H P P F A S V R N S E E A R T D R P S Q

AGTTGCGAGCTGAATGGCGAATGGGACGCGCCCTGTAGCGGCGCATTAAGCGCGGGGTGGTGGTTACGCGCAGCGTGACCGCTACACTTGCCAG 7900
TCAACGCGTCGGACTTACCGCTTACCTGCGCGGACATCGCGCGTAATTCGCGCGCCACACCAATGCGCGTCGCACTGGCGATGTGAACGGTC

D L R S L N G E V D A P C S G A L S A A G V V V T R S V T A T L A S

CGCCCTAGCGCCGCTCTTTTCGCTTTCTTCCCTTCTTCGCCACGTTGCGCGGCTTTCCTCGTAAGCTCTAAATCGGGGCTCCCTTTAGGGTTC 8000
GCGGGATCGCGGCGAGGAAAGCGAAAGGAAGGAAGGAGCGGTGCAAGCGGCGGAAAGGGGAGTTTCGAGATTAGCCCCGAGGGAAATCCCAAG

A L A P A P F A F F P S F L A T F A G F P R Q A L N R G L P L G F

CGATTAGTCTTACGGCACCTCGACCCCAAAAACTTGAATTAGGGTGATGGTTACGCTAGTGGCCATCGCCCTGATAGACGGTTTTTCGCCCTTTGA 8100
GCTAAATCAGGAAATGCGTGGAGCTGGGGTTTTTGAACCTAATCCACTACCAAGTGATCACCCTGAGCGGACTATCTGCCAAAAAGCGGAAACT

R F S A L R H L D P K K L D . G D G S R S G P S P . . T V F R P L

CGTTGGAGTCCAGTTCTTAAATAGTGGACTCTTGTTCCAACTGGAACAACACTCAACCTATCTCGGCTATTCTTTGATTATAAGGGATTTTGGC 8200
GCAACTCAGGTGCAAGAAATATCACCTGAGAACAAGGTTTGACCTTGTGTGAGTTGGGATAGAGCCAGATAAGAAAACCTAAATATCCCTAAACGG

T L E S T F F N S G L L F O T G T T L N P I S V Y S F D L . G I L P

GATTTCGGCTATTGGTTAAAAATGAGCTGATTTAACAATAATTAACGGAATTTTAAACAATAATTAACGCTTACAATTTAG 8285
CTAAAGCGGATAACCAATTTTTACTCGACTAAATTTTAAATTCGCTTAAATTTGTTTATAATTGCGAATGTTAAATC

I S A Y V L K N E L I . Q K F N A N F N K I L T L T I .

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Page 1

fig 55 pCB251 (1 > 8197) Site and Sequence

Enzymes : All 146 enzymes (No Filter)

Settings: Linear, Certain Sites Only, Standard Genetic Code

GACGGATCGGGAGATCTCCCGATCCCCTATGGTCGACTCTCAGTACAATCTGCTCTGATGCCGATAGTTAAGCCAGTATCTGCTCCCTGCTTGTGTGT
CTGCTTAGCCCTCTAGAGGGCTAGGGGATACCAGCTGAGAGTCATGTTAGACGAGACTACGGCGTATCAATTCGGTCATAGACGAGGGACGAACACACAA 100
T D R E I S R S P M V D S Q Y N L L . C R I V K P V S A P C L C V

GGAGGTCGCTGAGTAGTGCGCGAGCAAAATTAAGCTACAACAAGGCAAGGCTTGACCGACAATTGCATGAAGAATCTGCTTAGGGTTAGCGCTTTTGGC
CCTCCAGCGACTCATCAGCGCTCGTTTTAAATTCGATGTTGTTCCGTTCCGAACGGCTGTTAACGTACTTCTTAGACGAATCCCAATCCGCAAAACGC 200
G G R . V V R E Q N L S Y N K A R L D R O L H E E S A . G . A F C

CTGCTTCGCGATGTACGGGCAGATATACGCGTTGACATTGATTATTGACTAGTTATTAATAGTAATCAATTACGGGGTCATTAGTTCATAGCCCATATA
GACGAAGCGCTACATGCCCGGCTATATGCGCAACTGTAACATAAATGATCAATAATTATCATTAGTTAATGCCCGAGTAATCAAGTATCGGGTATAT 300
A A S R C T G Q I Y A L T L I I D . L L I V I N Y G V I S S . P I Y

TGGAGTTCGCGGTACATAACTTACGGTAATGGCCCGCCTGGCTGACCGCCCAACGACCCCGCCCATTGACGTCAATAATGACGTATGTTCCCATAGT
ACC TCAAGGCGCAATGATTGAATGCCATTACCGGGCGGACCGACTGGCGGGTTGCTGGGGGCGGGTAAC TGCAGTTATTACTGCATACAAGGGTATCA 400
G V P R Y I T Y G K V P A V L T A Q R P P P I D V N N D V C S H S

AACGCCAATAGGGACTTTCCATTGACGTCAATGGGTGGACTATTACGGTAACTGCCCACTTGGCAGTACATCAAGTGTATCATATGCCAAGTACGCC
TTGGGGTTATCCCTGAAAGGTAAGTGCAGTTACCCACCTGATAAATGCCATTTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTCATGCGGG 500
N A N R D F P L T S M G G L F T V N C P L G S T S S V S Y A K Y A

CCTATTGACGTCAATGACGGTAATGGCCCGCCTGGCATTATGCCAGTACATGACCTTATGGGACTTTCTCTACTTGGCAGTACATCTACGTATTAGTCA
GGATAACTGCAGTTACTGCCATTACCGGGCGGACCGTAATACGGGTGATGACTGGAATACCTGAAAGGATGAACCGTCATGTAGATGCATAATCAGT 600
P Y . R O . R . M A R L A L C P V H D L M G L S Y L A V H L R I S H

TCGCTATTACCATGGTGATGCGGTTTTGGCAGTACATCAATGGCGGTGGATAGCGGTTGACTCACGGGGATTTCGAAGTCTCCACCCCATTGACGTCAA
AGCGATAATGGTACCACCTACGCCAAAACCGTCATGTAGTTACCCGCACTATGCCAAACTGAGTGCCCTAAAGGTTGAGAGGTGGGGTAATGCGAGTT 700
R Y Y H G D A V L A V H O W A V I A V . L T G I S K S P P H . R Q

TGGGAGTTTGTGTTTGGCACCAAAATCAACGGGACTTTCCAAAATGTCGTAACAACCTCCGCCCATTGACGCAATGGGCGGTAGGCGGTGACGGTGGGAS
ACCTCAAAACAAACCGTGGTTTTAGTTGCCCTGAAAGGTTTTACAGCATGTTGAGGGGCGGTAAC TGC GTTTACCGGCCATCCGCACATSCCACCCTC 800
V E F V L A P K S T G L S K M S . O L R P I D A N G R . A C T V G

GTCTATATAAGCAGAGCTCTCTGGCTAACTAGAGAACCCTGCTTACTGGCTTATCGAAATTAATACGACTCACTATAGGGAGACCCAAGCTGGCTAGC
CAGATATATTGCTCTGAGAGACCGATTGATCTCTTGGGTGACGAATGACCGAATAGCTTTAATTATGCTGAGTGATATCCCTCTGGGTTGACCGGATCG 900
G L Y K Q S S L A N . R T H C L L A Y R N . Y D S L . G D P S V L A

GTTTAAACTTAAGCTTACCATGGGGGTTCTCATCATCATCATCATGGTATGGCTAGCATGACTGGTGGACAGCAATGGGTCGGGATCTGTACGAC
CAAATTTGAATTGGAATGGTACCCCCCAAGAGTAGTAGTAGTAGTAGTACCATACCGATCGTACTGACCACTGTGCTTTACCCAGCCCTAGACATGCTG 1000
F K L K L T M G G S H H H H H G M A S M T G G Q O M G R D L Y D

GATGACGATAAGGTACCCGGATCCTTCCGAGACCCACGACGATGTTACGGCTCAGTGCTGCTCCCTGGCTCCAGTGCTCTCCACCTACTCTCAG
CTACTGCTATTCCATGGGCTAGGAAGGCTCTGGGGTGCCTGCTACAAGTGCCGAGTCACGACAGGGACGGAGGTACGGAGGAGGTGGAAGAGGAGTCT 1100
D D D K V P G S F R D P T D D V H G S V L S L A S S A S S T V S S

T7 promoter priming site

Proband binding domain

pCB251 insert = U2

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fig 55 pCB251 (1 > 8197) Site and Sequence

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CTGAGGAGAGGATGCAATCTGAGCAAATCCGGAAGCTTCGTAGGGAAC TGGAAATCATCCAGGAAAAAGTGGCCACCTTGACGCTCAGCTTTCTGCCAA
GACTCCTCTCCTACGTTAGACTCGTTTAGGCCTTCGAAGCATCCCTTGACCTTAGTAGGGTCCTTTTCACCGGTGGAAC TGCAGAGTCGAAAGACGGT 120

pCB251 insert = U2

U2 ORF

A E E R M Q S E Q I R K L R R E L E S S Q E K V A T L T S Q L S A N

TGCTAATCTGGTGGCTGCTTTTGAGCAGAGCCTGGTGAATATGACATCCCGCTGCGACACCTGGCAGAGACGGCCGAGGAGAAGGACACTSAGCTGCTG
ACGATTAGACCACCGACGAAAAC TCGTCTCGGACCAC TTATACTGTAGGGCGGACGCTGTGGACCGTCTCTGCCGGCTCCTCTTCCTGTGACTCGACGAC 130

pCB251 insert = U2

U2 ORF

A N L V A A F E Q S L V N M T S R L R H L A E T A E E K D T E L L

GATTTGCGAGAAACCATAGACTTTCTGAAGAAAAAGAACTCTGAGGCCAGGCAGTCATTAGGGAGCCCTTAATGCCTCAGAAACACACCCAAAGAAC
CTAAACGCTCTTTGGTATCTGAAAGACTTCTTTTCTTGAGACTCCGGGTCGTCAGTAAGTCCCTCGGGAATTACGGAGTCTTTGGTGTGGGTTTCTTG 140

pCB251 insert = U2

U2 ORF

D L R E T I D F L K K K N S E A Q A V I O G A L N A S E T T P K E

TTCGGATCAAGAGACAAAACCTCTCAGATAGCATCTCAAGCCTCAACAGCATCACTAGCCATTCCAGCATCGGCAGCAGCAAGGATGCTGATGCGAAAAA
AAGCCTAGTTCTCTGTTTTGAGGAGTCTATCGTAGAGTTTCGGAGTTGTCGTAGTGATCGGTAAGGTCGTAGCCGTCGTCGTTCTACGACTACGCTTTTT 150

pCB251 insert = U2

U2 ORF

L R I K R O N S S O S I S S L N S I T S H S S I G S S K D A D A K I

GAAGAAAAAAGAGTTGGGTCTATGAGCTTCGAAGTTCCTTCAACAAAGCGTTCAGTATAAAAAAGGGGCCAAGTCAGCTTCTCATACTCGGATATA
CTTCTTTTTTTCTCAACCCAGATACTCGAAGCTTCAAGGAAGTTGTTTCGCAAGTCATATTTTCCCCGGGTTTCAGTCGAAGGAGTATGAGCCTATA 160

pCB251 insert = U2

U2 ORF

K K K K S V V Y E L R S S F N K A F S I K K G P K S A S S Y S D I

GAGGAGATTGCTACACCGACTCTTCAGCCCCCTCATCCCCAACTACAGCATGGTCTACAGAGACTGCTTACCCTCCATCAAGTCTCCACCTTGT
CTCCTCAACGATGTGGGCTGAGAAGTCGGGGAGTAGGGGTTTGATGTCGTACCAAGATGTCCTGACGAAGTGGGAGGTAGTTCAGGAGGTGGAACA 170

pCB251 insert = U2

U2 ORF

E E I A T P D S S A P S S P K L Q H G S T E T A S P S I K S S T L

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fig 55 pCB251 (1 > 8197) Site and Sequence

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CCTCCGTGGGCACTGATGTCACCGAGGGCCCTGCTCACCAGCCCCACACTAGGCTGTTCCATGCAAATGAGGAGGAGGCCAGAGAGAGAGAGGT
GGAGGCACCCGTGACTACAGTGGCTCCCGGGACGAGTGGGTCGGGGGGTGTGATCCGACAAGGTACGTTTACTCCTCCTCCTCGGTCTCTTCTCTCCA
pCB251 insert = U2
U2 ORF
S S V G T D V T E G P A H P A P H T R L F H A N E E E E P E K K E V
ATCGGAGCTGCGCTCTGAGCTATGGGAGAAGGAAATGAAGCTTACAGACATCCGCTTGGAGGCCCTCAACTCTGCCACCAACTGGATCAGCTTCGGGAG
TAGCCTCGACGCGAGACTCGATACCCCTTCTCTTACTTCAATGTCTGTAGGCGAACCTCCGGGAGTTGAGACGGGTGGTTGACCTAGTCGAAGCCCTC
pCB251 insert = U2
U2 ORF
S E L R S E L V E K E M K L T D I R L E A L N S A H Q L D Q L R E
ACCATGCAACATGCAGTTGGAGGTGGACCTGCTGAAAGCAGAGAATGACCGACTGAAGGTAGCCCCAGGCCCTCATCAGGCTCCACTCCAGGGCAGG
TGGTACGTGTTGTACGTCAACCTCCACC TGGACGACTTTCGTCTCTTACTGGCTGACTTCCATCGGGGTCCGGGGAGTAGTCCGAGGTGAGGTCCCGTCC
pCB251 insert = U2
U2 ORF
T M H N M O L E V D L L K A E N D R L K V A P G P S S G S T P G Q
TCCCTGGATCATCTGCATTATCTTCCCCACGCCGCTCCCTAGGCCTGGCACTACCCATTCTTTCGGCCCCAGTCTTGCAGACACAGACCTGTCACCCAT
AGGGACCTAGTAGACGTAATAGAAGGGGTGCGGCGAGGGATCCGGACCGTGAGTGGGTAAAGGAAGCCGGGGTCAGAACGTCGTGTC TGGACAGTGGGTG
pCB251 insert = U2
U2 ORF
V P G S S A L S S P R R S L G L A L T H S F G P S L A D T D L S P H
GGATGGCATCAGTACTTGTGGTCCAAAGGAGGAAGTGACCTCCGGGTGGTGGTGAGGATGCCCCGCAGCACATCATCAAAGGGGACTTGAAGCAGCAG
CCTACCGTAGTCATGAACACAGGTTTCTCTTCTACTGGGAGGCCACCACTCTTACGGGGGGTCTGTGTAGTAGTTTCCCTGAACCTTCGTCTGT
pCB251 insert = U2
U2 ORF
D G I S T C G P K E E V T L R V V V R M P P Q H I I K G D L K Q Q
GAATTCTTCTGGGCTGTAGCAAGGTCAAGTGGAAAAGTTGACTGGAAGATGCTGGATGAAGCTGTTTTCAAGTGTCAAGGACTATATTCTAAATGG
CTTAAGAAGGACCCGACATCGTTCAGTCACCTTTCAACTGACCTTC TACGACCTACTTCGACAAAAGGTTCAACAAGTTCTTGATATAAAGATTTTACC
pCB251 insert = U2
U2 ORF
E F F L G C S K V S G K V D V K M L D E A V F O V F K D Y I S K H
```

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fig 55 pCB251 (1 > 8197) Site and Sequence

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ACCCAGCCTCTACCTGGGACTAAGCACTGAGTCCATCCATGGCTACAGCATCAGCCACGTGAAACGAGTGTGGATGCAGAGCCCCGAGATGCCTCC
TGGGTCGGAGATGGGACCTGATTCGTGACTCAGGTAGGTACCGATGTCGTAGTCGGTGCACTTTGCTCACAACTACGTCTCGGGGGGCTCTACGGAGG 210

pCB251 insert = U2

U2 ORF

D P A S T L G L S T E S I H G Y S I S H V K R V L D A E P P E M P F

TGCCGTCGAGGTGTCATAACATATCAGTCTCCCTCAAAGGTCTGAAGGAGAAATGCGTCGACAGCCTGGTGTTCGAGACGCTGATCCCCAAGCCGATG
AACGGCAGCTCCACAGTTATTGTATAGTCAGAGGGAGTTTCAGAGCTTCTCTTACGCAGCTGTCGGACCACAAGCTCTCGGACTAGGGGTTCGGCTAC 220

pCB251 insert = U2

U2 ORF

C R R G V N N I S V S L K G L K E K C V D S L V F E T L I P K P H

ATGCAGCACTACATAAGCCTCCTGCTGAAGCACCGGCGCTCGTCTCTCGGGCCCCAGCGGCACGGGCAAGACCTACCTGACCAATCGCTTGGCCGAGT
TACGTCGTGATGATTTCGGAGGACGACTTCGTGGCCGCGGAGCAGGAGAGCCCGGGGTCGCCGTGCCCGTCTGGATGGACTGGTTAGCGAACC GGCTCA 230

pCB251 insert = U2

U2 ORF

M Q H Y I S L L L K H R R L V L S G P S G T G K T Y L T N R L A E

ACCTGGTGGAGCGCTCTGGCCGTGAGGTACAGAGGGCATCGTCAGCACCTTCAACATGCACCAGCAGTCTTGCAAGGATCTGCAACTGTATCTTTCCAA
TGGACCACCTCGCGAGACCGGCACTCCAGTGTCTCCCGTAGCAGTCGTGGAAGTTGTACGTGGTCGTCAGAACGTTCTTAGACGTTGACATAGAAAGGTT 240

pCB251 insert = U2

U2 ORF

Y L V E R S G R E V T E G I V S T F N M H Q Q S C K D L Q L Y L S N

CCTAGCCAACAGATAGACGGGAAACAGGAATTGGGGATGTGCCCTGGTGATTCTATTGGATGACCTGAGTGAAGCAGGCTCCATCAGTGAGTTGGTC
GGATCGGTTGGTCTATCTGGCCCTTTGTCTTAACCCCTACACGGGGACCACTAAGATAACCTACTGGACTCACTTCGTCGAGGTAGTCACTCAACCAAG 250

pCB251 insert = U2

U2 ORF

L A N Q I D R E T G I G D V P L V I L L D D L S E A G S I S E L V

AATGGGGCCCTCAGCTGCAAGTATCATAATGTCCCTATATTATAGGTACCACCAATCAGCCGTGAAAAATGACACCCAACCATGGCTTGCACTTGAGCT
TTACCCCGGGAGTGGACGTTTCATAGTATTACAGGGATATAATATCCATGGTGGTTAGTCGGACATTTTACTGTGGGTGGTACCGAACGTGAACTCGA 260

pCB251 insert = U2

U2 ORF

N G A L T C K Y H K C P Y I I G T T N Q P V K M T P N H G L H L S

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fig 55 pCB251 (1 > 8197) Site and Sequence

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TCAGGATGTTGACCTTC TCCAACAACGTGGAGCCAGCCAAATGGCTTCC TGGTTCGTTACCTGAGGAGGAAGCTGGTAGAGTCAGACAGCGACATCAATGC
AGTCC TACAAC TGAAGAGGT TGTTCACCTCGGTCGGT TACCGAAGGACCAAGCAATGGAC TCCTCCTTCGACCATCTCAGTCTGTCTGCTGTTACG 300

pCB251 insert = U2

U2 ORF

F R M L T F S N N V E P A N G F L V R Y L R R K L V E S D S D I N A

CAACAAGGAAGAGCTGCTTCGGGTGCTCGACTGGGTACCCAAGCTGTGGTATCATCTCCACACCTTCCTTGAGAAGCACAGCACCTCAGACTTCCTCATC
GTTGTTCTCTCGACGAAGCCCACGAGCTGACCCATGGGTTTCGACCATAGTAGAGGTGTGGAAGGAAC TCCTCGTGTCTGGAGTCTGAAGGAGTAG 310

pCB251 insert = U2

U2 ORF

N K E E L L R V L D V V P K L V Y H L H T F L E K H S T S O F L I

GGCCCTTGCTTCTTCTGTCTGTCCATTGGCATTGAGGACTTCCGGACCTGGTTCATTGACCTGTGGAACAACCTATCATTCCCTATCTACAGGAAG
CCGGGAACGAAGAAAGACAGCACAGGGTAACCGTAAC TCCTGAAGGCC TGGACCAAGTAAC TGGACACCTTGTGTAGATAGTAAGGGATAGATGTCCTTC 320

pCB251 insert = U2

U2 ORF

G P C F F L S C P I G I E D F R T V F I D L V N N S I I P Y L Q E

GAGCCAAGGATGGGATAAAGGTCCATGGACAGAAAGCTGCTTGGGAGGACCCAGTGGAATGGGTCCGGGACACACTTCCTTGCCATCAGCCCAACAAGA
CTCGGTTCTACCTATTTCAGGTACCTGTCTTTCGACGAACCTCTCGGTCACCTTACCCAGGCCCTGTGTGAAGGGACCGGTAGTCGGGTGTTCT 330

pCB251 insert = U2

U2 ORF

G A K D G I K V H G Q K A A V E D P V E V V R D T L P V P S A Q Q D

CCAATCAAGCTGTACCACCTGCCCCACCCACCGTGGGCCCTCACAGCATTGCCTCACCTCCCGAGGATAGGACAGTCAAAGACAGCACCCCAAGTTCT
GGTTAGTTTCGACATGGTGGACGGGGTGGGTGGCACCCGGGAGTGTCTAACGGAGTGGAGGGCTCTATCTGTCTAGTTTCGTCTGGGGTTCAAGA 340

pCB251 insert = U2

U2 ORF

O S K L Y H L P P P T V G P H S I A S P P E D R T V K D S T P S S

CTGGACTCAGATCCTCTGATGGCCATGCTGTGAAACTTCAAGAAGCTGCCAACTACATTGAGTCTCCAGATCGAGAAACCATCTGGACCCCAACCTTC
GACCTGAGTCTAGGAGACTACCGGTACGACGACTTGAAGTTCTTCGACGGTTGATGTAACTCAGAGGTCTAGCTCTTTGGTAGGACCTGGGGTTGGAAG 350

pCB251 insert = U2

U2 ORF

L D S D P L M A M L L K L Q E A A N Y I E S P D R E T I L D P N L

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fig 55 pCB251 (1 > 8197) Site and Sequence

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fig 55 pCB251 (1 > 8197) Site and Sequence

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GAATGTGTGTCAGTTAGGGTGTGAAAGTCCCCAGGCTCCCCAGGCAGGCAGAAGTATGCAAAGCATGCATCTCAATTAGTCAGCAACCAGGTGTGAAAA 560X
CTTACACACAGTCAATCCCACACCTTTCAGGGGTCCGAGGGGTCCGTCCGTCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTCCACACCTTT
N V C Q L G C G K S P G S P G R Q K Y A K H A S Q L V S N O V W I
GTCCCCAGGCTCCCCAGCAGGCAGAAGTATGCAAAGCATGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAATCCGCCCATCCGCCCTTAAT 570X
CAGGGGTCCGAGGGGTGCTCCGTCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTATCAGGGCGGGGATTGAGGCGGGTAGGGCGGGATTGA
V P R L P S R Q K Y A K H A S Q L V S N H S P A P N S A H P A P N
CCGCCAGTTCGCCCATCTCCGCCCATGGCTGACTAATTTTTTTTATTTATGCAAGGCGGAGGCGCCTCTGCCCTGAGCTATTCCAGAAGTAGT 580X
GGCGGGTCAAGGCGGGTAAGAGGCGGGTACCGACTGATTAATAAATAAATACGTCTCCGGCTCCGGCGGAGACGGAGACTCGATAAGGTCTTCATCA
S A Q F R P F S A P W L T N F F Y L C R G R G R L C L . A I P E V V
GAGGAGGCTTTTTTGAGGCGTAGGCTTTTGCAAAAAGCTCCCGGAGCTTGTATATCCATTTTCGGATCTGATCAAGAGACAGGATGAGGATCGTTTCG 590X
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R R L F W R P R L L Q K A P G S L Y I H F R I . S R D R M R I V S
CATGATTGAACAAGATGGATTGCACGCAGGTTCTCCGGCCGCTTGGGTGGAGAGGCTATTCGGCTATGACTGGGCACAACAGACAATCCGGCTGCTCTGAT 500X
GTACTAATCTGTTCTACCTAACGTGCGTCCAAGAGGCGGGCGAACCCACCTCTCCGATAAGCCGACTAGACCCGTGTTGCTGTTAGCCGACGAGACTA
H D . T R V I A R R F S G R L G G E A I R L . L G T T D N R L L .
GCCCGCGTGTCCGGCTGTGACGCGAGGGGCGCCCGGTTCTTTTGTCAAGACCGACCTGTCCGGTGCCCTGAATGAAGTGCAGGACGAGGCAGCGCGGC 510X
CGGCGGCACAAGGCCGACAGTCGCGTCCCCCGGGGCCAAGAAAACAGTTCTGGCTGGACAGGCCACGGGACTTACTTGACGTCTGCTCCGTGCGCGCG
C R R V P A V S A G A P G S F C O D R P V R C P E . T A G R G S A A
TATCGTGGCTGGCCACGACGGGCGTTCTTGGCGAGCTGTGCTCGACGTTGTACTGAAGCGGAAGGGACTGGCTGCTATTGGGCGAAGTGCCGGGGCA 520X
ATAGCACCGACCGGTGCTGCCCCGAAGGAACGCGTCGACACGAGCTGCAACAGTGACTTCGCCCTTCCC TGACCAGCATAACCCGCTTACGGCCCCGT
I V A G H D G R S L R S C A R R C H . S G K G L A A I G R S A G A
GGATCTCCTGTATCTACCTTGCTCCTCGCGAGAAAGTATCCATCATGGCTGATGCAATCGCGCGGCTGCATACGCTTGATCCGGCTACCTGCCATTTC 530X
CCTAGAGGACAGTAGAGTGAACGAGGACGGCTCTTTCATAGGTAGTACCGACTACGTTACGCCCGACGATATGCGAACTAGGCCGATGGACGGGTAAAG
G S P V I S P C S C R E S I H H G . C N A A A A Y A . S G Y L P I
GACCACCAAGCGAAACATCGCATCGAGCGAGCAGTACTCGGATGGAAGCCGGTCTTGTGATCAGGATGATCTGGACGAAGAGCATCAGGGGCTCGCGC 540X
CTGGTGGTTTCGCTTTGTAGCGTAGCTCGCTCGTGCATGAGCTTACCTCGGCCAGAACAGCTAGTCTTACTAGACCTGCTTCTCGTAGTCCCCGAGCGCG
R P P S E T S H R A S T Y S D G S R S C R S G . S G R R A S G A R A
CAGCCGAACGTGTCGCCAGGCTCAAGGCGCGCATGCCGACGGCGAGGATCTCGTCTGACCCATGGCGATGCCTGCTTGCCGAATATCATGTTGGAAAA 550X
GTCCGCTTGACAAGCGGTCCGAGTTCGCGCGCTACGGGCTGCCGCTCC TAGAGCAGCACTGGGTACCGCTACGGACGAACGGCTTATAGTACCACCTTTT
S R T V R O A Q G A H A R R R G S R R D P W R C L L A E Y H G G I
TGGCCGCTTTTCTGGATTATCAGCTGTGGCCGGCTGGGTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGAAGAGCTTGGC 560X
ACCGGCGAAAAGACCTAAGTAGCTGACACCGGCCGACCCACACCGCTGGCGATAGTCTGTATCGCAACCGATGGGCACATAAAGACTTCTCGAACCG
W P L F W I H R L W P A G C G G P L S G H S V G Y P . Y C . R A W
GGCGAATGGGTGACCGCTTCTCTGCTTTTACGGTATCGCGCTCCCGATTGCGAGCGCATCGCTTCTATCGCTTCTTGACGAGTTCTTCTGAGCGG 570X
CCGCTTACCGGACTGGCGAAGGAGCAGAAATGCCATAGCGGCGAGGGCTAAGCGTCCGCTAGCGGAAGATAGCGGAAGAACTGCTCAAGAGACTCGGC
R R M G . P L P R A L R Y R R S R F A A H R L L S P S . R V L L S G

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GACTCTGGGGTTCGAAATGACCGACCAAGCGACGCCAACCTGCCATCAGAGATTTCGATTCCACCGCCGCCCTTCTATGAAAGGTTGGGCTTCGGATC
CTGAGACCCCAAGCTTTACTGGCTGGTTTCGCTGCGGGTTGGACGGTAGTGCTCTAAAGCTAAGGTGGCGGCGGAAGATACCTTCCAACCGAAGCCTTAG
T L G F E M T D Q A T P N L P S R D F D S T A A F Y E R L G F G I 580X

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CAAAAGGCCCTGCGGCCGACCTACTAGGAGGTGCGGCCCTAGAGTAGCAGCTCAAGAAGCGGGTGGGGTTGAACAAATAACGTGCAATATTACCAATGT
V F R D A G V M I L Q R G D L M L E F F A H P N L F I A A Y N G Y 590X

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TTATTTCTGTTATCGTAGTGTTTAAAGTGTTTATTTCTGAAAAAAGTGACGTAAGATCAACACCAACAGGTTTGAGTAGTTACATAGAATAGTACAGAC
K . S N S I T N F T N K A F F S L H S S C G L S K L I N V S Y H V C 600X

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ATATGGCAGCTGGAGATCGATCTCGAACCGCATTAGTACCAGTATCGACAAGGACACACTTTAACAATAGGCGAGTGTTAAGGTGTGTTGATGCTCGG
I P S T S S . S L A . S V S . L F P V . N C Y P L T I P H N I R A 610X

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CCTTCGTATTTACATTTTCGACCCACGGATTACTCACTCGATTGAGTGTAATTAACGCAACGCGAGTGACGGGCGAAAGGTCAGCCCTTTGGACAGCA
G S I K C K A V G A . . V S . L T L I A L R S L P A F Q S G N L S 620X

GCCAGCTGCATTAATGAATCGGCCAACGCGGGGAGAGGCGGTTTGGCTATTGGGCGCTCTCCGCTTCTCGCTCACTGACTCGCTGCGCTCGGTCGT
CGGTCGACGTAATTACTTAGCCGGTTGCGCGCCCTCTCCGCCAACGCATAACCCGCGAGAAGGCGAAGGAGCGAGTGACTGAGCGACGCGAGCCAGCA
C Q L H . . I G Q R A G R G G L R I G R S S A S S L T O S L R S V V 630X

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AGCCGACGCGCTCGCCATAGTCGAGTGAGTTTCCGCCATTATGCCAATAGGTGCTTAGTCCCTATTGCGTCTTCTTGTACACTCGTTTTCCGGTC
R L R R A V S A H S K A V I R L S T E S G D N A G K N M . A K G Q 640X

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Q K A R N R K K A A L L A F F H R L R P P D E H H K N R R S S Q R 650X

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CCGCTTTGGGCTGTCTGATTTCTATGGTCCGCAAGGGGGGACCTTCGAGGGAGCACGCGAGAGGACAAGGCTGGGACGGCGAATGGCCTATGGACAG
V R N P T G L . R Y Q A F P P G S S L V R S P V P T L P L T G Y L S 660X

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GCGGAAAGAGGGAAGCCCTTCGCACCGGAAAGAGTTACGAGTGCGACATCCATAGAGTCAAGCCACATCCAGCAAGCGAGGTTCGACCCGACACAGTG
A F L P S G S V A L S O C S R C R Y L S S V . V V R S K L G C V H 670X

GAACCCCCGTTTCAGCCGACCGCTGCGCCTTATCCSGTAACATATGCTTGTAGTCAACCCGGTAAGACACGACTTATCGCCACTGGCAGCAGCCACTG
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E P P V Q P D R C A L S G N Y R L E S N P V R H D L S P L A A A T 680X

GTAACAGGATTAGCAGAGCGAGGTATGTAGGCGGTGCTACAGAGTTCTTGAAGTGGTGGCTTAACACGGCTACACTAGAAGGACAGTATTTGGTATCTG
CATTTGCTCTAATCGTCTCGCTCCATACATCCGCCACGATGCTCAAGAATTCACCACGGATTGATGCCGATGTGATCTTCTGTGCTAACCATAGAC
G N R I S R A R Y V G G A T E F L K V V P N Y G Y T R R T V F S I C 690X

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CGCTCTGCTGAAGCCAGTTACCTTCGGAAAAAGAGTTGGTAGCTCTTGATCCGGCAACAAACCACCGCTGGTAGCGGTGGTTTTTTTGTGTTGCAAGCAG
GCGAGACGACTTCGGTCAATGGAAGCCTTTTTCTCAACCATCGAGAAC TAGGCCGTTTGTGGTGGCGACCATCGCCACCAAAAAACAAACGTTTCGTCT
A L L K P V T F G K R V G S S . S G K Q T T A G S G G F F V C K Q 750

CAGATTACGCGCAGAAAAAGGATCTCAAGAAGATCCTTTGATCTTTCTACGGGGTCTGACGCTCAGTGAACGAAAACTCACGTTAAGGGATTTTGG
GTCTAATGCGCGCTTTTTTCTAGAGTTCTCTAGGAACTAGAAAAAGTGGCCAGACTGCGAGTCACCTTGCTTTTGAGTGCAATCCCTAAAAACC
Q I T R R K K G S Q E D P L I F S T G S D A Q W N E N S R . G I L 760

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V M R L S K R I F T . I L L N . K . S F K S I . S I Y E . T W S D S 770

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Y Q C L I S E A P I S A I C L F R S S I V A . L P V V . I T T I R 780

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E G L P S G P S A A M I P R D P R S P A P D L S A I N O P A G R A 790

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E R R S G P A T L S A S I Q S I N C C R E A R V S S S P V N S L R N 800

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GCAACAACGGTAACGATGTCGTAGCACCACAGTGCAGCAGCAACCATACCGAAGTAAGTCGAGGCCAAGGGTTCGTAGTTCGCTCAATGATCTAGG
V V A I A T G I V V S R S S F G M A S F S S G S Q R S R R V T . S 810

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P M L C K K A V S S F G P P I V V R S K L A A V L S L M V M A A L 820

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H N S L T V M P S V R C F S V T G E Y S T K S F . E . C M R R P S C 830

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S C P A S I R D N T A P H S R T L K V L I I G K R S S G R K L S R 840

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I L P L L R S S S M . P T R A P N . S S A S F T F T S V S G . A K 850

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GTCTTCCGTTTACGGCGTTTTTCCCTTATCCCGCTGTGCTTTTACAACCTATGAGTATGAGAAGGAAAAAGTTATAATAAC TTCGTAAATAGTCCC
T G R O N A A K K G I R A T R K C . I L I L F L F O Y Y . S I Y O G 860

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. TTATTGTCATGAGCGGATACATATTTGAATGTATTAGAAAAATAACAAATAGGGGTTCCGCGCACATTTCCCGAAAAGTGCCACCTGACGTC
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Y C L M S G Y I F E C I . K N K Q I G V P R T F P R K V P P D V

Figure 56

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fig 56 pNP10

Page 1

10 20 30 40 50 60 70
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360 370 380 390 400 410 420
CAGCAACCGCTTCAGCAACTAACCCAAATTCCAACTTTCCACAAATGTCAACATCCAGGCCTCAGACTCC 420
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710 720 730 740 750 760 770
TCATACTGGAAATATGATCTGAAACAAATTTGTGCTATTCTCAAAATGTTTAAAGCATGTTTGAAGAT 770
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1060 1070 1080 1090 1100 1110 1120
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1410 1420 1430 1440 1450 1460 1470
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1760 1770 1780 1790 1800 1810 1820
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2110 2120 2130 2140 2150 2160 2170
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CTTGCCTGCTTCACTCTAATCCAGACATGATTTCCAAAGAGAGTTTCAATTTTAAATGTTTCCAGTTTCT 2450

Figure 5b

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fig 5b pNP10

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2460 2470 2480 2490 2500 2510 2520
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GATTCATCAAAAGATGGTATCAAGCCAAAGACGTCTGGACGAAACCACCCCATCATCAACCACGCAI 2590
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CACATCTGCCAAGAGCTTAGGTATCCGATCCTTCCGGCTTCTTTTAGAAATGAGAGGATGAGAGGCA 2730
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2810 2820 2830 2840 2850 2860 2870
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3160 3170 3180 3190 3200 3210 3220
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3860 3870 3880 3890 3900 3910 3920
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4560 4570 4580 4590 4600 4610 4620
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lig pNP10

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CAGTGC TGGCATAACCAIGAGTGAFAACACTGCGGCCAACTTACTTCTGACAACGATCGGAGGACCGAAG 5600
5610 5620 5630 5640 5650 5660 5670
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5960 5970 5980 5990 6000 6010 6020
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CTGAGCGTCAGACCCGTCAGAAAAGATCAAAGGATCTTCTTGAGATCTCTTTTTCGCGCGTAATCTGC 6230
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6310 6320 6330 6340 6350 6360 6370
TTCGGAAGGTAACTGGCTTCAACGAGCGCGAGATACCAAAATACGTCTTCTAGTGTAGCGGTAGTTAGG 6370
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GCCAGTGGCGATAAGTCTGTCTTACCGGGT TGGAC TCAAGACGATAGTTACCGGATAAGGCGGACGGGT 6510
CGGGCTGAACCGGGGGTCTGTGCACACAGCCAGCTTGGAGCGAACGACCTACACCGAACTGAGATACCT 6580
ACAGCGTGAGCATTGAGAAAGCGCCAGCTTCCCGAAGGGAGAAAGGCGGACAGGTATCCGGTAAGCGCG 6650
6660 6670 6680 6690 6700 6710 6720
AGGGTCGGAAACAGGAGAGCGCACGAGGAGCTTCCAGGGGAAACGCC TGGTAICTTTATAGTCTGTCTC 6720
GGTTTCGCCACCTCTGACTTGAGCGTCTGATTTTGTGTATGCTCGTCAGGGGGGCGGAGCCTATGGAAATA 6790
CGCCAGCAACCGCGGCTTTTACGGTTCTTGGCTTTTTC TGGCTTTTGTCTCAATGTTCTTTCTCTCG 6860
TTATCCCTGATCTGTGGAFAACCGTATTACCGCTT TGGAGTGAGCTGATACCGCTCGCCGAGCGGAA 6930
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7010 7020 7030 7040 7050 7060 7070
GGCTTGGCGATTCATTAATGCAGCTGGACGACAGCTTTCCCGAG TGGAAAGCGGCGACTGACCGCAAC 7070
GCAATTAATGTGAGATTAAGTCACTCATTAGGCAACCCAGGCTTTACACTTTATGCTTCCCGCTCGTATGT 7140
TGTTTGAATTGTGAGCGGAFAACAATTTACACAGGAAGACAGTATGACCATGAT TACGCCAAGCTGTA 7210
AGTTTAAACATGATCTTACFAACFAACTATTCTCATTTTAAATTTTACAGAGCTTAAATATGGCTGAAATCA 7280
CTCACAACGATGGATACCTAACAATTTGGAAATGAAAT 7319

Figure 57

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10 20 30 40 50 60 70
ATACCAAGATACGCAAGCTTGCATGCCTGCAGGAATTCGATATCAAGCTTATCGATACCGTCGACCT 70
AGAGGATCAGAAAGAAATTGGACCAACTAGCCACATCCAATTAAGCCACCGCGGTTTCTAAGTGAGTTTAA 140
TTTTGAGTTTACGACACAAAAATGTGTTCTTTAATAACTATCTTCGACTTGAGTCATTCTGTATGACT 210
AGTTGTTGAGTGATTTTTCATTGAGAAAAATATTAAGGAACAATTTACTTTTGCATTTTGCCTTAA 280
TTTGATTAGTTTTCGATCAACTAGATCTTACAAAACTTGCAATACAATTCATTTCAGATTACCCCTC 350
360 370 380 390 400 410 420
GCCACGTGTCGCCACGTACGCAACCGGTTTCAGCAACTAACCCAAAATTCCAACTTTCACAAAAATGCAACA 420
TCCAGGCTTCAGACTCCACAGTCAAGAATATCGAAAAATTTGTAAGAAATTTTATTTTGGAGCTCAAACTTGT 490
ATAAAAATGCCAGAAAAAGAAGATGATAAAAAATGTAGTTTTCGCAAAAACTTCCACCTTTATTGCTCTAA 560
TATGACGGCTTATATCTCAATTTTCTTGAATTTATCAAAAAATTTTCACATATACAAAATGAGAAAAAGT 630
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710 720 730 740 750 760 770
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1410 1420 1430 1440 1450 1460 1470
ATTTATTTTCATATTTAAATTTTGTGCAATTAGGGATAAACACGACTTTTAAAAAGTTTATTTAAAAAAACG 1470
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1760 1770 1780 1790 1800 1810 1820
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2110 2120 2130 2140 2150 2160 2170
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GTTAAGCCATGATTGACTTCTTGTGCTGAGTCTAATCCAGACTAGATTTCCAAGAGATTTTCAATTTT 2450

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2810 2820 2830 2840 2850 2860 2870
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3160 3170 3180 3190 3200 3210 3220
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3510 3520 3530 3540 3550 3560 3570
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3860 3870 3880 3890 3900 3910 3920
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4210 4220 4230 4240 4250 4260 4270
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4560 4570 4580 4590 4600 4610 4620
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TTGAGCAAAAGCTTAGAAAACTCACTCAACACATGATCGATCCAACTTGAAGCTTGAAGAGGCAATACG 4900

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TCCGATCTCAAGGAAC TCTTGACAACATTGATGTGATTGAGTTGAAGCAAGAGCTCAAAGAACGCGATA 5250
5260 5270 5280 5290 5300 5310 5320
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GTACATCAGCTAGTCAATCTTCGAAACGATCTCTGGCTGCAACTCAATCAAGGTTACTGTAAACGTGGA 5530
CA TCGCTGGAGAAATCAGTTGATCGT TAAACGGGACTTGAAGCAGCAGGAATTCTTCTGGGCTGTAGC 5600
5610 5620 5630 5640 5650 5660 5670
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5960 5970 5980 5990 6000 6010 6020
GACCTACCTGACCAATCGCTTGGCCGAGTACC TGGTGGAGCGCTCTGGCCGTGAGGTACAGAGGGCATC 6020
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6310 6320 6330 6340 6350 6360 6370
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CATCAATGCCAACAAAGGAGAGCTGCT TCGGGTGTCTGACTGGGTACCCAAGCTGTGGTATCATCTCCAC 6440
AGCTTCTTGAAGAGCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 6510
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6660 6670 6680 6690 6700 6710 6720
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7010 7020 7030 7040 7050 7060 7070
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CTAGGAAACGAA TGGTGGGGTGGGCT TGGGAACCTGTGCCCC TAAACACATTTACTGGCTTCTTAAT 7140
GACT TGGGGAAGAGATGATCT TGGTCTT TCCCTTGAC TCTTGGTTCAATTACAAC TCTGGCTCTT 7210
CTGGGGAGGGGTTCAGAAAATCATCAAAACACTGCAAGCAGTTCCCCGGAATTCAGCTTGGAC TTAACAGG 7280
CTGAAC TTGCTCAAAAGAGCCGAATTCCAGCACTGACCTGCCCTCCCCATGGTAT TGA TATCTGAGCTCCGC 7350

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7360 7370 7380 7390 7400 7410 7420
ATCGGGCGGCTGTCATCAGATCGCCATCTCGCGCCCGTGCCCTCGACTTCFAAGGCCAACTACGCTTCAAC 7420
ATCCCTACATGCTCTTTCGCCGIGIGGCCACCCCTATTTTTGTTATTATCAAAAAAACCTCTTCTTA 7430
ATTTCTTTTGTGTTTGTAGCTTCTTTTAAAGTCACCTCTAACAATGAAATTGTGTAGATTCAAAAAATAGAATT 7440
AATTCGTAATAAAAAAGTCGAAAAAAATTTGTGCTCCCTCCCCCTTAATAATAATTCTATCCCAAAATCT 7450
ACACAAATGTCTGTGTACACITCTTAAGTGTTTTTTACTTCTGATAAAATTTTTTTGAAACATCATAGAA 7460

7710 7720 7730 7740 7750 7760 7770
AAAACCGCACACAAAAATACCTTATCATATGTTACGTTTCAGTTTATGACCGCAATTTTATTTCTTCGCA 7770
CGTCTGGGCTCTCATGACGTCAAATCATGCTCATCGTGAAAAAATTTTGGAGTATTTTGGAAATTTTTC 7780
AATCAAGTGAAAGTTTATGAAATTAATTTTCTGCTTTTGTCTTTTGGGGGTTTCCCTATTTGTTGTCA 7790
AGAGTTTCGAGGACGGCGTTTCTTTGCTAAATCACAAAGTATTGATGAGCAGGATCAAGAAAGATCGG 7800
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8060 8070 8080 8090 8100 8110 8120
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AGACGGTCACAGCTTGTCTGTAAGCGGATGCGGGGAGCAGACAAGCCCGTCAGGGCSCGTCAGCGGGTGT 8140
TGGCGGGTGTGCGGGGCTGGCTTAACATTCGGCATCAGAGCAGATTGTACGAGAGTCACCAATATCCGG 8150
TGTGAAATACCGCACAGATGCGTAAGGAAGAAAATACCGCATCAGGCGGCCCTGAAGGCCCTGCTGATAC 8160

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CGCTTTTTTGGCGCATTTTGCCTTCTGTTTTTGTCTACCCAGAAACGCTGGTGAAGTAAAAGATGCTG 8500
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8760 8770 8780 8790 8800 8810 8820
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9110 9120 9130 9140 9150 9160 9170
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9460 9470 9480 9490 9500 9510 9520
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9810 9820 9830 9840 9850 9860 9870
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 10160 10170 10180 10190 10200 10210 10220
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 10510 10520 10530 10540 10550 10560 10570
 ACTCATTAGGCACCCACGGCTTACAGTTTATGCTTCCGGCTCGTATGTTGTGTGGAAATGAGCGGAG 10570
 AACAAATTCACACAGGAAACAGCT 10594

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Fig 58

Page

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10 20 30 40 50 60 70
ATGACCATGATTACGCCAAGCTTGCATGCCGTCAGGTCGACTctagAAATGFAAACCTGTCATTTCTGTG 70
TATTTTCAGCACCGCAGAGAGCACCATAAACAGCTACAACAAGAGCACGAACGTCGGCTCCAACAGAAAA 140
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360 370 380 390 400 410 420
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710 720 730 740 750 760 770
CTGTGCGTGGTGCAGAGCTCATGAATAAGTGATGGGGCACCGAGATGTCGACTCCCTCCCATTAATCTCTT 770
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1060 1070 1080 1090 1100 1110 1120
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1410 1420 1430 1440 1450 1460 1470
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1760 1770 1780 1790 1800 1810 1820
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GAAGACTATTGAGCAAAACAAACAGCCCAATAACAGCGGTGTTGGTGGTGGTGGTGGTGGTGGTGGTGGT 1960
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pLM3 (1 > 10847) Site and Sequence

Enzymes :

100 of 146 enzymes (Filtered)

Settings:

Linear, Certain Sites Only, Standard Genetic Code

(Seq 361)

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G G R . V V R E Q N L S Y N K A R L D R Q L H E E S A . G . A F C
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TT promoter priming site

ProBond binding domain

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pLM3 (1 > 10847) Site and Sequence

Page 2

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ORF pLM1
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ORF pLM1
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ORF pLM1
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ORF pLM1
S D S D L M G K T M T E D D D I T T G V D E S S S I S S G L S D A

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pLM3 (1 > 10847) Site and Sequence

Page 3

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insert pLM1

ORF pLM1

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insert pLM1

ORF pLM1

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insert pLM1

ORF pLM1

L K M E P G T S K W R R E R P E S C D D S S K G G E L K K P I S L

GGCCACCCTGGTTCCTTGAAGAAGGGCAAGACCCACCTGTGGCTGTAACCTCCCCATCACTCACACAGCCCAGAGTGCCCTCAAAGTCGCAGGCAAAAC
CCGGTGGGACCAAGGGACTTCTTCCCGTCTTGGGGTGGACACCGACATTGAAGGGGGTAGTGAGTGTGTCGGGTCTACGGGAGTTTCAGCGTCCGTTTG

insert pLM1

ORF pLM1

G H P G S L K K G K T P P V A V T S P I T H T A Q S A L K V A G F

CTGAGGGCAAAGCTACAGACAAGGGTAAGCTTGCACTGAAGAATACTGGGCTCCAACGCTCTCTCTGATGCTGGTGGGACCGCTGAGTGATGCTAA
GACTCCCGTTTCGATGCTGTTCCTTCAAGCTCACTTCTTATGACCCGAGGTGCGAGGAGGAGACTACGACCAAGCCCTGGCGGACTCACTACGAT

insert pLM1

ORF pLM1

P E G K A T O K G K L A V K N T G L Q R S S S D A G R D R L S D A F

GAAGCCCCCTCGGGCATTGCTCGCCCTCCACTTCGGGATCCTTCGGCTACAAGAAGCTCTCTCTGCCACAGGCACAGCCACTGTATGCAAACTGGT
CTTCGGGGGAGCCGTAACGAGCGGGGAGGTGAAGCCCTAGGAAGCCGATGTTCTTCGGAGGAGGACGGTGTCCGTGTCGGTGACAGTACGTTTGACCA

insert pLM1

ORF pLM1

K P P S G I A R P S T S G S F G Y K K P P P A T G T A T V M Q T G

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GGTTCAGCCACTCTCAGCAAGATCCAGAAGTCCTCAGGCATCCCTGTCAAGCCAGTAAATGGGCGCAAGACTAGCTTAGATGTTTCCAACAGCGCAGAGC
CCAAGTCGGTGAGAGTCGTTCTAGGTCTTCAGGAGTCGGTAGGGACAGTTCCGGTCATTTACCCGCGTTCTGATCGAATCTACAAAGGTTGTCGCGTCTCG
2300
-----insert pLM1-----
-----ORF pLM1-----
G S A T L S K I O K S S G I P V K P V N G R K T S L D V S N S A E
CAGGATTCTGGCTCCGAGAGCCGTTCTAACATCCAGTACCGCAGCCTGCCCCGGCCAGCCAAGTCAAGTTCATGAGCGTGACCGGCGGGGCGGGTGG
GTCTTAAGGACCGAGGACCTCGGGCAAGATTGTAGGTTCATGGCGTCGGACGGGGCCGGTCGGTTCAGTTCAAGATACTCGCACTGGCCGCCCGCCCCACC
2400
-----insert pLM1-----
-----ORF pLM1-----
P G F L A P G A R S N I O Y R S L P R P A K S S S M S V T G G R G G
ACCTCGCCCTGTGAGCAGCAGCATTGACCCAGTCTCCTCAGCACCAAGCAGGGAGGCCTTACGCCCTTCAGACTGAAGGAGCCTACCAAGGTAGCCAGT
TGGAGCGGGACACTCGTCGCTGTAACGGGGTCAGAGGAGTCGTGGTTCGTCCCTCCGGAATGCGGAAGGTCTGACTTCCTCGGATGGTTCCATCGGTCA
2500
-----insert pLM1-----
-----ORF pLM1-----
P R P V S S S I D P S L L S T K Q G G L T P S R L K E P T K V A S
GGGCGGACCACTCCAGCCCTGTCAATCAGACAGATCGGGAAAAGGAGAAGGCCAAAGCCAAGGCAGTGGCCTTGGACTCAGACAACATCTCCTTGAAGA
CCCGCCTGGTGAGGTGCGGGACAGTTAGTCTGTC TAGCCCTTTTCCCTCTCCGGTTTCGGTTCGGTCACCGGAACCTGAGTCTGTTGTAGAGGAACTTCT
2600
-----insert pLM1-----
-----ORF pLM1-----
G R T T P A P V N Q T D R E K E K A K A K A V A L D S D N I S L)
GTATTGGCTCCCCAGAGAGTACTCCAAGAACCAAGCAAGCCACCCACAGCCACCAAGCTGGCAGAGCTGCCACCAACCCCTCTCAGGGCCACAGGGAA
CATAACCGAGGGGTCTCTCATGAGGGTCTTGGTTCGTTCCGTGGGTGTCGGTGGTTCGACCGTCTCGACGGTGGTTGGGGAGAGTCCCGGTGTCCT
2700
-----insert pLM1-----
-----ORF pLM1-----
S I G S P E S T P K N Q A S H P T A T K L A E L P P T P L R A T A)
GAGCTTTGTCAAACCACTCTACATAGCCAATCTTGACAAGGTCAACTCCAACAGTCTGGATCTACCATCATCCAGTGATACCACCATGCTTCAAAGGTG
CTCGAAACAGTTTGGTGGGAGTGATCGGTTAGAACGTTCAGTTGAGGTGTGACAGCTAGATGGTAGGTAGGTCACTATGGTGGGTACGAAGTTTCCAG
2800
-----insert pLM1-----
-----ORF pLM1-----
S F V K P P S L A N L D K V H S N S L D L P S S S D T T H A S I V

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CCAGATCTGCATGCTACAAGCTCAGCATCTGGGGGCCCTCTCCCTTCTGCTTACCCCCAGTCCGGCACCCATCCTCAATATTAACTCAGCCAGCTTCT
GGTCTAGACGTACGATGTTTCGAGTCGTAGACCCCCGGGAGAGGGAAGGACGAAGTGGGGTTCAGGCCGTGGGTAGGAGTTATAATTGAGTCGGTCAAGGA
2900
-----insert pLM1-----
-----ORF pLM1-----
P D L H A T S S A S G G P L P S C F T P S P A P I L N I N S A S F
CCAGGGCCTGGAGCTAATGAGTGGTTTCAGTGTGCCAAAAGAGACCCGCATGTACCCCAAACTCTCAGGCCTGCACAGGAGCATGGAGTCCCTCCAGAT
GGTCCCGGACCTCGATTACTCACCAAAGTCACACGGTTTCTCTGGGCGTACATGGGGTTTGAGAGTCCGGACGTGTCTCGTACCTCAGGGAGGTCCTA
3000
-----insert pLM1-----
-----ORF pLM1-----
S Q G L E L M S G F S V P K E T R M Y P K L S G L H R S M E S L Q M
GCCAATGAGCCTCCCCAGTGCCTTCCCCAGCAGTACTCCCGTCCCCACCCACCTGCTCCCCCTGCTGCTCCACAGAAGAAGAGACGGAAGAGCTGACT
CGGTTACTCGGAGGGGTCACGGAAGGGGTCGTATGAGGGCAGGGGTGGGGTGACGAGGGGGACGACGAGGGTGTCTTCTTCTGCTTCTCGACTGA
3100
-----insert pLM1-----
-----ORF pLM1-----
P M S L P S A F P S S T P V P T P P A P P A A P T E E E T E E L T
TGGAGTGGGAAGCCCCAGAGCTGGGCAAC TGGACAGTAATCAGCGGGATCGGAACACTCTTCCCAAGAAAGGGCTCAGGTACCAGCTTCAGTCCAGGAGG
ACCTCACCTTCGGGGTCTCGACCCGTTGACCTGTCTATTAGTCGCCCTAGCCCTGTGAGAAGGGTCTTTCCCGAGTCCATGGTCGAAGTCAGGGTCCCTC
3200
-----insert pLM1-----
-----ORF pLM1-----
V S G S P R A G Q L D S N Q R D R N T L P K K G L R Y Q L Q S Q E
AGACCAAGGAGAGGCGACATTCCCATACCATTTGGTGGGC TGCCTGAATCCGATGACCAGTCAGAGCTGCCTTCTCCCCCTGCACCTTCCCATGCTCTGAG
CTGGTTCTCTCCGCTGTAAAGGTATGGTAACCAACCCGACGGACTTAGGCTACTGGTCAGTCTCGACGGAAGAGGGGGACGTGAAGGGTACAGAGACTC
3300
-----insert pLM1-----
-----ORF pLM1-----
E T A E R R H S H T I G G L P E S D D Q S E L P S P P A L P M S L
TGC AAAGGCCAAC TTACCAACATAGTGAGTCCCACTGCGGCCACCAACGCCAAGAATCACCCGCTCCAAACAGCATCCCCACCCACGAGGGCGGCTTCGAG
ACGTTTCCCGGTGAATGGTTGTATCACTCAGGGTGACGCCGGTGGTGGGTTCTTAGTGGGCGAGGTTGTCTAGGGGTGGGTGCTCCGCCGGAAGCTC
3400
-----insert pLM1-----
-----ORF pLM1-----
A K G Q L T N I V S P T A A T T P R I T R S N S I P T H E A A F E

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CTGTACAGCGGCTCCCAAAATGGGGAGCACCTGTCCCTGGCCGAGAGACCCAAGGGAATGATTCGGTCAGGATCCTTCCGAGACCCACGGACGATGTTT 351
GACATGTGCGGAGGGTTTACCCCTCGTGGGACAGGGACCGGCTCTCTGGGTTCCCTTACTAAGCCAGTCTTAGGAAGGCTCTGGGGTGCCGTGCTACAAAG
-----insert pLM1-----
-----ORF pLM1-----
L Y S G S Q M G S T L S L A E R P K G M I R S G S F R D P T D D V
ACGGCTCAGTGCTGTCCCTGGCCGCCAGTGCCCTCCACCTACTCTCAGCTGAGGAGAGGATGCAATCTGAGCAAAATCCGGAAGCTTCGTAGGGAAC 360
TGCCGAGTCACGACAGGGACCGAGGTCACGGAGGAGGTGGATGAGGAGTCGACTCCTCTCCCTACGTTAGACTCGTTTAGGCCCTCGAAGCATCCCTTGA
-----insert pLM1-----
-----ORF pLM1-----
H G S V L S L A S S A S S T Y S S A E E R M Q S E Q I R K L R R E L
GGAATCATCCAGGAAAAAGTGCCACCCTTGACGTCTCAGCTTTCTGCCAATGCTAATCTGGTGGCTGCTTTTGAGCAGAGCCTGGTGAATATGACATCC 370
CCTTAGTAGGGTCCCTTTTACCCTGGAACTGCAGAGTCGAAAGACGGTTACGATTAGACCACCGACGAAAACCTCGTCTCGGACCACTTATACGTAGG
-----insert pLM1-----
-----ORF pLM1-----
E S S Q E K V A T L T S Q L S A N A N L V A A F E Q S L V N M T S
CGCCTGCSACACTGGCAGAGACGGCCGAGGAGAAGGACACTGAGCTGCTGGATTTGCGAGAAACCATAGACTTTCTGAAGAAAAAGAACTCTGAGGCC 380
GCGGACGCTGTGGACCGTCTCTGCCGGCTCCTCTTCCGTGACTCGACGACCTAAACGCTCTTTGGTATCTGAAAGACTTCTTTTCTTGAGACTCCGGS
-----insert pLM1-----
-----ORF pLM1-----
R L R H L A E T A E E K D T E L L D L R E T I D F L K K K N S E A
AUGCACTCATTGAGGGAGCCCTTAATGCCTCAGAAACCAACCCAAAGAACCTCGGATCAAGAGACAAACCTCCTCAGATAGCATCTCAAGCCTCAACAG 390
TCCGTCACTAAGTCCCTCGGGAATTACGGAGTCTTTGGTGTGGGTTTCTTGAAGCCTAGTTCCTGTTTGGAGGAGTCTATCGTAGAGTTCGGAGTTGTC
-----insert pLM1-----
-----ORF pLM1-----
Q A V I Q G A L N A S E T T P K E L R I K R Q N S S D S I S S L N S
CATCACTAGCCATTCAGCATCGGCAGCAGCAAGGATGCTGATGCGAAAAAGAAGAAAAAAGAGTTGGGTCTATGAGCTTCGAAGTTCTTCAACAAA 400
GTAGTGATCGGTAAAGGTCGTAGCCGTCGTCGTTCCTACGACTACGCTTTTCTCTTTTCTCAACCCAGATACTCGAAGCTTCAAGGAAGTTGTT
-----insert pLM1-----
-----ORF pLM1-----
I T S H S S I G S S K D A D A K K K K K K S V V Y E L R S S F H I
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GCGTTCAGTATAAAAAAGGGGCCAAGTCAGCTTCCTCATAC TCGGATATAGAGGAGATTGCTACACCCGACTCTTCAGCCCCCTCATCCCCAACTAC
CGCAAGTCATATTTTTCCTCCGGGTTTCAGTCGAAGGAGTATGAGCCTATATCTCTCTAACGATGTGGGCTGAGAAGTCGGGGGAGTAGGGGTTGATG 410

----- insert pLM1 -----

----- ORF pLM1 -----

A F S I K K G P K S A S S Y S D I E E I A T P D S S A P S S P K L

AGCATGGTTCCACAGAGACTGCTTCACCTCCATCAAGTCCTCCACCTTGCTCTCCGTGGGCACTGATGTCACCGAGGGCCC TGCTCACCCAGCCCCCA
TCGTACCAAGGTGCTCTGACGAAGTGGGAGGTAGTTTCAGGAGGTGAACAGGAGGCACCCGTGACTACAGTGGCTCCCGGACGAGTGGGTCGGGGGGT 420

----- insert pLM1 -----

----- ORF pLM1 -----

Q H G S T E T A S P S I K S S T L S S V G T D V T E G P A H P A P H

CACTAGGCTGTTCCATGCAAAATGAGGAGGAGGACCCAGAGAAGAAGGAGGTATCGGAGCTGCGCTCTGAGCTATGGGAGAAGGAAATGAAGCTTACAGAC
GTGATCCGACAAGGTACGTTTACTCCTCCTCGGTCTCTTCTCTCCATAGCCTCGACGCGAGACTCGATACCCTCTCTCTTACTTCGAATGTCTG 430

----- insert pLM1 -----

----- ORF pLM1 -----

T R L F H A N E E E E P E K K E V S E L R S E L V E K E M K L T D

ATCCGCTTGGAGGGCCCTCAACTCTGCCACCAACTGGATCAGCTTCGGGAGACCATGCACAACTGCAGTTGGAGGTGGACCTGCTGAAAGCAGAGAATG
TAGGCGAACCTCCGGGAGTTGAGACGGGTGGTTGACCTAGTCAAGCCCTCTGGTACGTGTTGTACGTCAACCTCCACCTGGACGACTTTCGTCCTTAC 440

----- insert pLM1 -----

----- ORF pLM1 -----

I R L E A L N S A H Q L D Q L R E T M H N M O L E V D L L K A E N

ACCGACTSAAGGTAGCCCCAGGCCCTCATCAGGCTCCACTCCAGGGCAGGTCCTTGATCATCTGCATTATCTTCCCCACGCCGCTCCCTAGGCTGGC
TGGCTGACTTCCATCGGGGTCCGGGAGTAGTCCGAGGTGAGGTCCGTCCTAGGGACCTAGTAGACGTAAATAGAAGGGGTGCGGCGAGGGATCCGGACCC 450

----- insert pLM1 -----

----- ORF pLM1 -----

D R L K V A P G P S S G S T P G Q V P G S S A L S S P R R S L G L A

ACTCACCATTCCTTCGGCCCCAGTCTTGACAGACAGACCTGTCAACCATGGATGGCATCAGTACTTGTGGTCCAAGGAGGAAGTGACCTCCGGGTS
TGAGTGGGTAAAGGAAGCCGGGGTCAGAACGTCGTGTCTGGACAGTGGGTACCTACCGTAGTCATGAACACCAAGGTTTCTCTTCACTGGGAGGCCAC 460

----- insert pLM1 -----

----- ORF pLM1 -----

L T H S F G P S L A D T D L S P M D G I S T C G P K E E V T L R V

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GTGGTGAGGATGCCCCCGCAGCACATCATCAAAGGGGACTTGAAGCAGCA3GAATTCTTCCTGGGCTGTAGCAAGGTCAGTGGAAAAGTTGACTGGAAGA
CACCACCTCTACGGGGGCGTCGTGTAGTAGTTTCCCTGAAC TTCGTCGTCCTTAAGAAGGACCCGACATCGTTCAGTCACCTTTTCAACTGACCTTCT 4700

insert pLM1

ORF pLM1

V V R H P P Q H I I K G O L K Q Q E F F L G C S K V S G K V D V K

TGCTGGATGAAGCTGTTTTCCAAGTGTTCAGGACTATATTTCTAAATGGACCCAGCCTCTACCTGGGACTAAGCACTGAGTCCATCCATGGCTACAG
ACGACCTACTTCGACAAAAGGTTCAAGTTCCTGATATAAAGATTTTACCTGGGTCGGAGATGGGACCCCTGATTCGTGACTCAGGTAGGTACCGATGTC 4800

insert pLM1

ORF pLM1

M L D E A V F Q V F K D Y I S K M D P A S T L G L S T E S I H G Y S

CATCAGCCACGTGAACAGAGTGTGGATGCAGAGCCCCCGAGATGCC TCTTGGCGTCGAGGTGTCAATAACATATCAGTCTCCCTCAAAGGTCCTGAAG
GTAGTCGGTGCACCTTGTCTACAACCTACGTC TCGGGGGGCTCTACGGAGGAACGGCAGC TCCACAGTTATTGTATAGTCAGAGGGAGTTTCCAGACTTC 4900

insert pLM1

ORF pLM1

I S H V K R V L D A E P P E M P P C R R G V N N I S V S L K G L I

GAGAAATGCGTCGACAGCC TGGTGTTCGAGACGCTGATCCCCAAGCCGATGATGCAGCACTACATAAGCCTCCTGCTGAAGCACC6GCGCTCGTCTCT
CTCTTTACGCAGCTGTGCGACCAAGCTCTGCGACTAGGGGTTGCGCTACTACGTCGTGATGTATTGCGAGGACGACTTCGTGGCCGCGGAGCAGSAGA 5000

insert pLM1

ORF pLM1

E K C V D S L V F E T L I P K P M M Q H Y I S L L L K H R R L V L

CGGGCCCCAGCGGCACGGGCAAGACCTACCTGACCAATCGCTTGGCCGAGTACCTGGTGSAGCGCTCTGGCCGTGAGGTCACAGAGGGCATCGTCAGCAE
GCCCCGGGTCGCGGTGCGCGTCTGGATGGACTGGTTAGCGAACC6GCTCATGGACCACCTCGCGAGACCGGCACCTCCAGTGCTCTCCCGTAGCAGTCTG 5100

insert pLM1

ORF pLM1

S G P S G T G K T Y L T N R L A E Y L V E R S G R E V T E G I V S T

CTTCAACATGCACCAGCAGTCTTTGCAAGGATCTGCAACTGTATCTTTCCAACCTAGCCAACAGATAGACCGGGAAACAGGAATTGGGGATGTGCCCTG
GAAGTTGTACGTGGTCGTCAGAAGCTTCCTAGACGTTGACATAGAAAGGTTGATCGGTTGGTCTATCTGGCCCTTTGTCCTTAACCCCTACACGGGGAC 5200

insert pLM1

ORF pLM1

F N M H O Q S C K D L O L Y L S N L A N O I D R E T G I G D V F L

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GTGATTCTATTGGATGACCTGAGTGAAGCAGGCTCCATCAGTGAGTTGGTCAATGGGGCCCTCACCTGCAAGTATCATAAATGTCCTATATTATAAGTA
CAC TAAGATAACCTACTGGACTCACTTCGTCCGAGGTAGTCACTCAACCAGTTACCCGGGAGTGGACGTTTCATAGTATTTACAGGGATATAATATCCAT
5300
-----insert pLM1-----
-----ORF pLM1-----
V I L L D D L S E A G S I S E L V N G A L T C K Y H K C P Y I I G
CCACCAATCAGCCTGTAAAAATGACACCCAACCATGGCTTGCACTTGAGCTTCAGGATGTGACCTTCTCCAACAACGTGGAGCCAGCCAATGGCTTCCT
GGTGGTTAGTCGGACATTTTACTGTGGGTTGGTACCGAAGCTGAACTCGAAGTCCTACAAC TGAAGAGGTTGTTGCACCTCGGTCGGTTACCGAAGGA
5400
-----insert pLM1-----
-----ORF pLM1-----
T T N Q P V K M T P N H G L H L S F R M L T F S N N V E P A N G F L
GGTTCGTACCTGAGGAGGAAGCTGGTAGAGTCAGACAGCGACATCAATGCCAACAGGAAGAGCTGCTTCGGGTGCTCGACTGGGTACCCAAGCTGTGG
CCAAGCAATGGACTCCTCCTTCGACCATCTCAGTCTGTCGCTGTAGTTACGGTTGTTCTCTCGACGAAGCCACAGAGCTGACCCATGGGTTCGACACC
5500
-----insert pLM1-----
-----ORF pLM1-----
V R Y L R R K L V E S D S D I N A N K E E L L R V L D V V P K L V
TATCATCTCCACACCTTCCTTGAGAAGCACAGCACCTCAGACTTCCTCATCGGCCCTTGCTTCTTTCTGTCGTGCCATTGGCATTGAGGACTTCGGGA
ATAGTAGAGGTGTGGAAGGAACCTTCGTGTGCTGGAGTCTGAAGGAGTAGCCGGGAACGAAGAAAGACAGCACAGGGTAACCGTAACCTCTGAAGGCT
5600
-----insert pLM1-----
-----ORF pLM1-----
Y H L H T F L E K H S T S D F L I G P C F F L S C P I G I E D F R
CCTGGTTTCATTGACCTGTGGAACAACCTATCATTCCTTCATCTACAGGAAGGAGCCAAGGATGGGATAAAGGTCCATGGACAGAAAGCTGCTTGSGAGGA
GGACCAAGTAAC TGGACACCTTGTTGAGATAGTAAGGGATAGATGTCTTCCTCGGTTCTTACCCTATTTCCAGGTACCTGTCTTCGACGAACCTCCT
5700
-----insert pLM1-----
-----ORF pLM1-----
T V F I D L V N N S I I P Y L Q E G A K D G I K V H G Q K A A W E D
CCCAGTGGAATGGGTCGGGACACACTTCCTGSCCATCAGCCCAACAAGACCAATCAAAGCTGTACCACCTGCCCCACCCACCGTGGGCCCTCACAGC
GGGTACCTTACCCAGGCCCTGTGTGAAGGGACCGGTAGTCGGGTTGTTCTGGTTAGTTTCGACATGGTGGACGGGGTGGGTGGCACCCGGGAGTGTCTG
5800
-----insert pLM1-----
-----ORF pLM1-----
P V E V V R D T L P V P S A Q Q D Q S K L Y H L P P P T V G P H S
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ATTGCCTCACCTCCCGAGGATAGGACAGTCAAAGACAGCACCCCAAGTCTCTGGACTCAGATCCTCTGATGGCCATGCTGCTGAAACTTCAAGAAGCTE
TAA CGGAGTGGAGGGCTCCATCCTGTCACTTCTGTCTGGGGTCAAGAGACCTGAGTCTAGGAGACTACCGGTACGACGACITTGAACTCTTCGAC 530X

— insert pLM1 —

— ORF pLM1 —

I A S P P E D R T V K D S T P S S L D S D P L M A M L L K L Q E A

CCAAC TACAT T GAGTCTCCAGATCGAGAAACCATCCTGGACCCCAACCTTCAGGCAACACTTTAAGGGTTCGGCAATCACTGTACCCCCGGACAGCAGA
GGTGTAGTAAC T CAGAGGCTAGCTCTTTGGTAGGACCTGGGGTGGAAAGTCCGTGTGAAATCCCAAGCCGTTAGTGACAGTGGGGGCTGTCTGCT 540X

— insert pLM1 —

— ORF pLM1 —

A N Y I E S P D R E T I L D P N L Q A T L . G F G N H C H P R T A E

ACGCTGGCATCAGCTATCTTAGCTCCTCTCTCCCTCTCTCTTTCAGAGCACTGGCTCTCCAGCCCCAGGAGGAGAACAGGAGGGAGGAGAGATGAA
TGCGACCGTAGTCGATAGAATCGAGGAGGAGAGGGGAGAGGAGAAAGTCTCGTGACCGAGAGGTCGGGGTCTCTCTTGTCTCCCTCTCTCTACTT 550X

— insert pLM1 —

R V H Q L S . L L L S P L L F Q S T G S P A P G G E Q E G G G D E

AGAGGAGGGACAGGTTCTTGGTGCTGTACCTTTGAGAAC TTCCTAGGAAGGAATGGTGGGTGGCGTTTGGGAAC TTGTGCCCC TAAACACATTTACTG
TCTCTCTCTGTCCAAGAACCACGACATGGAAC TCTTGAAGGATCCTTCTTACCACCCACC GCAACCC TTGAACACGGGGATTGTGTAAATGAC 560X

— insert pLM1 —

R G G T G S V C C T F E N F L G R N G G V A F G N L C P L N T F T

GCCTCTCTAATGACTTTGGGAAAAGATGATTCTGGGTCTTTCCCTTGACTTCTTGTTCATTACAAAC TCCTGGGCTTTCTGGGGAGGSGTTCAGAA
CGGAGGAGATTACTGAAACCCCTTTTCTACTAAGACCCAGAAAGGGAAC TGAAGAACAAGT TAATGTTTGAGGACCCGAAAGACCCCTCCCAAGCTT 570X

— insert pLM1 —

G L L . . L V G K D D S G S F P . L L V S I T N S W A F V G G V Q

AACATCAAAACACTGCAGCAGTTCCTCCGGAATTCAGCTTGGACTTAACCAGGCTGAAC TTGCTCAAAAGAAGCCGAATTCAGCACACTGGCGGCTTAA
TTGTAGTTTTGTGACGTCGTAAGGGGCTTAAGTCGAACCTGAATTGGTCCGACTTGAACGAGTTTCTTCGGCTTAAGGTCGTGTGACCGCGGCAA 580X

— insert pLM1 —

T S K H C S S S P E F S L D L T R L N L L K R S R I P A H W R P L

CTAGTTCTAGAGGGCCGTTTAAACCCGCTGATCAGCCTCGACTGTGCCTTCTAGTTGCCAGCCATCTGTTGTTTGGCCCC TCCCCGTCCTTCTTGAC
GATCAAGATCTCCCGGCAAAATTTGGGCGACTAGTCGGAGCTGACACGGAAGATCAACGGTCGGTAGACAACAACGGGGAGGGGGACGGAAGGAAGCTE 590X

— insert pLM1 —

L V L E G P F K P A D Q P R L C L L V A S H L L F A P P P C L P .

CCTGGAAGGTGCCACTCCCACTGTCTTTCTAATAAAATGAGGAAATGTCATCGCATTTGCTGAGTAGGTGTCATTCTATTCTGGGGGGTGGGTTGGGG
GGACCTTCCACGGTGAGGGTGACAGGAAAGGATTATTTACTCTTTAAGTAGCGTAACAGACTCATCCACAGTAAGATAAGACCCCCACCCACCCG
P V K V P L P L S F P N K M R K L H R I V . V G V I L F V G V S V G 600X

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CAGGACAGCAAGGGGAGGATTGGGAAGACAATAGCAGGCATGCTGGGGATGCGGTGGGCTCTATGGCTTC TGAGGCGGAAAGAACCAAGCTGGGGCTCTA
GTCCTGTCGTTCCCTCTCTAACCTTCTGTTATCGTCCGTACGACCCCTACGCCACCCGAGATACCGAAGAC TCCGCTTTCTTGGTCGACCCCGAGAT 6700
R T A R G R I G K T I A G M L G M R V A L V L L R R K E P A G A L

GGGGGTATCCCCACGCGCCTGTAGCGGCGCAATTAAGCGCGGGGGTGTGGTGGTTACGCGCAGCGTGACCGCTACACTTGCCAGCGCCCTAGCGCCCGC
CCCCATAGGGGTGCGCGGACATCGCCGCTAATTGCGCGCGCCACACCACCAATGCGCGTCGCACTGGCGATGTGAACGGTCGCGGGATCGCGGGCG 6800
G G I P T R P V A A H . A R R V W V L R A A . P L H L P A P . R P

TCCTTTCGCTTTCTCCCTTCCTTTCGCGCACGTTGCGCGGCTTCCCGTCAAGCTCTAAATCGGGGCATCCCTTTAGGGTTCCGATTAGTGCTTTA
AGGAAAGCGAAAGAGGGAAGAGCGGTGCAAGCGCGCGAAAGGGGCAGTTGAGATT TAGCCCCGTAGGGAATCCCAAGGCTAAATCACGAAAT 6900
L L S L S S L P F S P R S P A F P V K L . I G A S L . G S D L V L Y

CGGCACCTCGACCCCAAAAACTTGATTAGGGTGATGGTTCACGTAGTGGGCCATCGCCCTGATAGACGGTTTTTCGCCCTTTGACGTTGGAGTCCACGT
GCGGTGGAGCTGGGGTTTTTGAAC TAATCCCACTACCAAGTGCATCACCGGTAGCGGGACTATCTGCCAAAAAGCGGGAACGCAACCTCAGGTGCA 7000
G T S T P K N L I R V M V H V V G H R P D R R F F A L . R V S P R

TCTTTAATAGTGGACTCTTGTTCACAACTGGAACAACAC TCAACCTATCTCGGTCTATTCTTTTGAATTATAAGGGATTTTGGGGATTTCGGCCTATTG
AGAAATTATCACCTGAGAACAGGTTTGACCTTGTGTGAGTTGGGATAGAGCCAGATAAGAAACTAAATATTCCTAAAACCCCTAAAGCGGATAAC 7100
S L I V D S C S K L E Q H S T L S R S I L L I Y K G F V G F R P I

GTTAAAAATGAGCTGATTTAACAAAAATTAACGCGAATTAATCTGTGGAATGTGTGTCAGTTAGGGTGTTGAAAGTCCCCAGGCTCCCCAGGCAGGC
CAATTTTTTACTCGACTAAATTGTTTTAAATTGCGCTTAATTAAGACACCTTACACACAGTCAATCCACACCTTTTCAGGGGTCCGAGGGGTCCGTCGC
G . K M S . F N K N L T R I N S V E C V S V R V W K V P R L P R Q A 7200

AGAAGTATGCAAGCATGCATCTCAATTAGTCAGCAACCAGGTGTGGAAGTCCCCAGGC TCCCCAGCAGGCAGAAGTATGCAAGCATGCATCTCAATT
TCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTCCACACCTTTTCAGGGGTCCGAGGGGTGTCGCTCTTCATACGTTTCGTACGTAGAGTTAA 7300
E V C K A C I S I S Q O P G V E S P Q A P Q Q A E V C K A C I S I

AGTCAGCAACCATAGTCCCGCCCTAAC TCCGCCATCCCGCCCTAAC TCCGCCAGTTCCGCCCATTCGCCCCCATGGCTGACTAATTTTTTTTAT
TCAGTCGTTGGTATCAGGGCGGGGATTGAGGCGGGTAGGGCGGGGATTGAGGCGGGTCAAGGCGGGTAAGAGGCGGGTACC GACTGATTAATAAAAAATA 7400
S O O P . S R P . L R P S R P . L R P V P P I L R P M A D . F F L

TTATGCAGAGGCGGAGGCGCCTCTGCC TCTGAGCTATTCCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCTTAGGCTTTTGCAAAAAGCTCCCGGAGCT
AATACGTCTCCGGCTCCGGCGGAGACGGAGACTCGATAAGGTCTTCATCACCTCTCCGAAAAACCTCCGGATCCGAAAACGTTTTTCGAGGGCCCTCGA 7500
F M Q R P R P P L P L S Y S R S S E E A F L E A . A F A K S S R E L

TGTATATCCATTTTCGGATCTGATCAAGAGACAGGATGAGGATCGTTTCGCATGATTGAACAAGATGGATTGCACGCAGGTTCTCCGGCCGCTTGGGTGS
ACATATAGGTAAGGCTAGACTAGTTCTCTGTCTTACTCTAGCAAGCGTACTAATCTGTCTACCTAACGTGCGTCCAAGAGGCGGCGAACCACCC 7600
V Y P F S D L I K R Q D E D R F A . L N K M D C T O V L R P L G W

AGAGGCTATTTCGGCTATGACTGGGCAACAAGACAATCGGCTGCTCTGATGCCGCGTGTTCGGCTGTGACGCGAGGGGCGCCGGTCTTTTTTGTCAA
TCTCCGATAAGCCGATGACTGACCGTGTGTCTGTAGGCGACGAGACTACGGCGGCAAGGCGGACAGTCGCGTCCCCGCGGGCCAAGAAAAACGTT 7700
R G Y S A M T G H N R O S A A L M P P C S G C Q R R G A R F F L S

GACCGACCTGTCCGGTCCCTGAATGAAC TGCAGGACGAGGACGCGCGGCTATCGTGGCTGGCCACGACGCGCGTTCCTTGCGCAGCTGTGCTCGACGTT
CTGGCTGGACAGGCCACGGGACTTACTTGACGTCTGCTCCGTCGCGCGGATAGCACCGACCGGTGCTGCCCGCAAGGAACGCGTCGACACGAGCTGCAA 7800
R P T C P V P . M N C R T R O R G Y R G V P R R A F L A Q L C S T L

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GTCAC TGAAGCGGGAAGGGACTGGCTGCTATTGGGCGAAGTGCCGGGCGAGGATCTCCTGTCATCTCACCTTGCTCCTGCCGAGAAAGTATCCATCATGG
CAGTGACTTCGCCCTTCCTGACCGACGATAACCCGCTTCACGGCCCGCTCCTAGAGGACAGTAGAGTGGAACGAGGACGGCTCTTTCATAGGTAGTACC 7900
S L K R E G T G C Y W A K C R G R I S C H L T L L L P R K Y P S V

CTGATGCAATGCGGCGGCTGCATACGCTTGATCCGGCTACCTGCCCATTCGACCACCAAGCGAAACATCGCATCGAGCGAGCAGTACTCGGATGGAAAGC
GACTACGTTACGCCCGCAGCTATGCGAACTAGGCCGATGGACGGGTAAGCTGGTGGTTCGCTTTGTAGCGTAGCTCGCTCGTGCATGAGCCTACCTTCG 8000
L M Q C G G C I R L I R L P A H S T Y K R N I A S S E H V L G W K

CGGTCTTGTGCGATCAGGATGATCTGGACGAAGAGCATCAGGGGCTCGCGCCAGCCGAACCTGTTCCGCCAGGCTCAAGGCGCGCATGCCCGACGGCGAGGAT
GCCAGAACAGCTAGTCTACTAGACCTGCTTCTCGTAGTCCCCGAGCGGGTCGGCTTGACAAGCGGTCCGAGTTCGCGCGTACGGGCTGCCGCTCTTA 8100
P V L S I R M I W T K S I R G S R Q P N C S P G S R R A C P T A R I

CTCGTCGTGACCCATGGCGATGCCTGCTTGCCGAATATCATGGTGGAAAAATGGCCGCTTTTCTGGATTTCGACTGTGGCCGGCTGGGTGTGGCGGACC
GAGCAGCACTGGGTACCGCTACGGACGAACGGCTTATAGTACCACCTTTTACCGCGAAAAGACCTAAGTAGCTGACACCGGCCGACCCACACCGCTGG 8200
S S . P M A M P A C R I S W V K M A A F L D S S T V A G W V V R T

GCTATCAGGACATAGCGTTGGCTACCGTGATATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCGCTTCTCGTGCTTTACGGTATCGCCGCTCCCGA
CGATAGTCTGTATCGCAACCGATGGGCACTATAACGACTTCTCGAACC CGCGCTTACCGGACTGGCGAAGGAGCAGCAAATGCCATAGCGGCGAGGGCT 8300
A I R T . R W L P V I L L K S L A A N G L T A S S C F T V S P L P

TTGCGAGCGCATCGCTTCTATCGCTTCTTGACGAGTCTTCTGAGCGGGACTCTGGGGTTCGAAATGACCGACCAAGCGACGCCCAACCTGCCATCAC
AAGCGTCGCTAGCGGAAGATAGCGGAAGAAGTGC TCAAGAAGACTCGCCCTGAGACCCCAAGCTTTACTGGCTGGTTCGCTCGGGTTGGACGGTAGTG 8400
I R S A S P S I A F L T S S S E R D S G V R N D R P S D A Q P A I T

GAGATTTGATTCACCGCGCCTTCTATGAAAGTTGGGCTTCGGAATCGTTTTCCGGGACGCCGGCTGGATGATCTCCAGCGCGGGGATCTCATGCT
CTCTAAAGCTAAGGTGGCGGCGGAAGATACTTTCAACCCGAAGCCTTAGCAAAAGGCCCTGCGGCGGACCTACTAGGAGGTGCGGCCCTTAGAGTACGA 8500
R F R F H R R L L . K V G L R N R F P G R R L D O P P A R G S H A

GGAGTTCTTCCGCCACCCCAACTGTTTTATTGCAGCTTATAATGGTTACAATAAAGCAATAGCATCACAAATTCACAAATAAAGCATTTTTTTCAGTS
CCTCAAGAAGCGGGTGGGGTTGAACAAATAACGTGCAATATTACCAATGTTTATTTCGTTATCGTAGTGTTTAAAGTGTTTATTCGTAAGAAAAAGTGAC 8600
S V L R P P O L V Y C S L . W L Q I K O . H H K F H K . S ! F F T

CATTCTAGTTGTGGTTTGTCCAAATCATCAATGTATCTTATCATGCTGTATACCGTCGACCTCTAGCTAGAGCTTGGCGTAATCATGGTCATAGCTGT
GTAAGATCAACACCAACAGGTTTGTAGTAGTTACATAGAATAGTACAGACATATGGCAGCTGGAGATCGATCTCGAACCAGCATTAGTACCAGTATCGACA 8700
A F . L W F V Q T H Q C I L S C L Y T V D L . L E L G V I M V I A V

TTCTGTGTGAAATGTTATCCGCTCACAATCCACACAACATACGAGCGGAAGCATAAAGTGTAAGCCTGGGGTGCTTAATGAGTGAGCTAAC TCAC
AAGGACACACTTTAACAATAGGCGAGTGTTAAGGTGTGTGTATGCTCGGCCCTCGTATTTCACATTTCCGAGCCCCAGGATTACTCACTCGATTGAGTG 8800
S C V K L L S A H N S T Q H T S R K H K V . S L G C L M S E L T H

ATTAATTGCGTTGCGCTCACTGCCCGCTTTCCAGTCGGGAAACCTGTCGTGCCAGCTGCATTAATGAATCGGCCAACGCGCGGGGAGAGSGGGTTTTCGT
TAATTAAACGCAACGCGAGTGACGGGCGAAAGGTCAGCCCTTTGGACAGCAGGTCGACGTAAATTACTTAGCCGGTTGCGCGCCCTCTCCGCCAAACGCA 8900
I N C V A L T A R F P V G K P V V P A A L M N R P T R G E R R F A

ATTGGGCGCTCTTCGCTTCTCGCTCACTGACTCGCTGCGCTCGGTCTGTTCGGCTGCGGCGAGCGGTATCAGCTCACTCAAAGGCGGTAAACAGGTTAT
TAACCCGCGAGAAGGCGAAGGAGCGAGTGACTGAGCGACGCGAGCCAGCAAGCCGACGCCGCTCGCCATAGTCGAGTGAGTTTCGCCATTATGCCAATA 9000
V A L F R F L A H . L A A L G R S A A A S G I S S L K G G N T V I

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CCACAGAATCAGGGGATAACGCAGGAAGAACATGTGAGCAAAAGGCCAGCAAAAGGCCAGGAACCGTAAAAAGGCCGCTTGCTGGCGTTTTTCCATAG
GGTGTCTTAGTCCCTATTGCGTCTTTCTTTGACACTCGTTTTCCGGTCGTTTTCCGGTCTTGGCATTTTTCCGGCGCAACGACCGCAAAAAGGTATC 910
H R I R G . R R K E H V S K R P A K G Q E P . K G R V A G V F P .
GCTCCGCCCCCTGACGAGCATCACAAAAATCGACGCTCAAGTCAGAGGTGGCGAAACCCGACAGGACTATAAGATACCAGGCGTTTCCCTTGGAAAGC
CGAGCGGGGGGACTGCTCGTAGTGTTTTAGCTGCGAGTTCAGTCTCCACCGCTTTGGCTGTCTGATATTTCTATGGTCCGCAAGGGGGGACCTTCG 920
A P P P . R A S Q K S T L K S E V A K P D R T I K I P G V S P V K
TCCCTCGTGGCTCTCCTGTTCCGACCTGCCGCTTACCGGATACCTGTCCGCTTTCTCCCTTCGGGAAGCGTGGCGTTTTCTCAATGCTACGCTGTA
AGGGAGCACGCGAGAGGACAAGGCTGGGACGGCAATGGCTATGGACAGCGGAAAGAGGGAAGCCCTTCGACCGCGAAAGAGTTACGAGTGGCAGAT 930
L P R A L S C S D P A A Y R I P V R L S P F G K R G A F S M L T L
GGTATCTCAGTTCGGTGTAGGTCGTTGCTCCAAGCTGGGCTGTGTGCACGAACCCCGTTAGCGCCGACCGCTGCGCTTATCCGGTAACATCGTCT
CCATAGAGTCAAGCCACATCCAGCAAGCGAGGTTGACCCGACACAGTGCTTGGGGGGCAAGTCGGGCTGGCGACGCGGAATAGGCCATTGATAGCAGA 940
V S Q F G V G R S L Q A G L C A R T P R S A R P L R L I R . L S S
TGAGTCCAACCCGTAAGACACGACTTATGCCACTGGCAGCAGCCACTGGTAACAGGATTAGCAGAGCGAGGTATGTAGGCGGTGCTACAGAGTTCTTG
ACTCAGGTTGGGCCATTCTGTCTGAATAGCGGTGACCGTCGCTGGTGACCATGTCTTAATCGTCTCGCTCCATACATCCGCCACGATGTCTCAAGAAC 950
V Q P G K T R L I A T G S S H V . Q D . Q S E V C R R C Y R V L
AAGTGGTGGCTAACTACGGCTACACTAGAGGACAGTATTTGGTATCTGCGCTCTGCTGAAGCCAGTTACCTTCGGAAAAAGAGTTGGTAGCTCTTGAT
TTCACCACCGGATTGATGCCGATGTGATCTTCTGTCTATAAACCATAGACGCGAGACGACTTCGGTCAATGGAAGCCCTTTTCTCAACCATCGAGAATA 960
E V V A . L R L H . K D S I V Y L R S A E A S Y L R K K S W . L L I
CCGGCAACAAACCACCGCTGGTAGCGGTGGTTTTTTGTTTGAAGCAGCAGATTACGCGCAGAAAAAAGGATCTCAAGAAGATCCTTTGATCTTTTC
GGCCGTTTTGTTGGTGGCGACCATGCCACCAAAAAAACAAAGTTCGTCGCTCAATGCGGCTTTTTTTCTTAGAGTTCTCTAGGAACTAGAAAAAG 970
R O T N H R V . R W F F C L Q A A D Y A O K K R I S R R S F D L F
TACGGGGTCTGACGCTCAGTGAACGAAACCTACGTTAAGGGATTTTGGTCATGAGATTATCAAAAAGGATCTTACCTAGATCCTTTTAAATTAATAA
ATGCCCCAGACTGCGAGTCACCTTGCTTTTGAAGTCAATTCCTAAACAGTACTCTAATAGTTTTCTTAGAAGTGGATCTAGGAAAAATTAATTTTT 980
Y G V . R S V E R K L T L R D F G H E I I K K D L H L D P F K L I
TGAAGTTTTAAATCAATCTAAAGTATATATGAGTAAACTTGGTCTGACAGTTACCAATGCTTAATCAGTGAGGCACCTATCTCAGCGATCTGTCTATTTC
ACTTCAAAATTTAGTTAGATTTCATATATACTCATTTGAACCAGACTGTCAATGGTTACGAATTAGTCACTCCGTGGATAGAGTCGCTAGACAGATAAAG 990
M K F . I N L K Y I . V N L V . O L P M L N Q . G T Y L S D L S I S
GTTATCCATAGTTGCCTGACTCCCCGTCGTGTAGATAACTACGATACGGGAGGGCTTACCATCTGGCCCCAGTGCTGCAATGATACCGGAGACCCACG
CAAGTAGGTATCAACGGACTGAGGGGCGAGCATCTATTGATGCTATGCCCTCCCGAATGGTAGACCGGGTCACGACGTTACTATGGCGCTCTGGGTGC 1000
F I H S C L T P R R V D N Y D T G G L T I W P Q C C N D T A R P T
CTCACCGGCTCCAGATTTATCAGCAATAAACGACGCCGGAAGGGCGGAGCGCAGAAGTGGTCTGCAACTTTATCCGCCCTCCATCCAGTCTATTAAT
GAGTGGCCGAGGTCTAAATAGTCGTTATTTGGTGGTCCGCTTCCCGGCTCGCGTCTTACCAGGACGTTGAAATAGGCGGAGGTAGGTGAGATAATTA 1010
L T G S R F I S N K P A S R K G R A O K V S C N F I R L H P V Y
TGTTCGCCGGAAGCTAGAGTAAGTAGTTGCCAGTTAATAGTTTGCACAACGTTGTTGCCATTGCTACAGGCATCGTGGTGTACGCTCGTCTTGGTA
ACAACGGCCCTTCGATCTCATTCATCAAGCGGTCAATTAACAACGCGTTGCAACAACGGTAACGATGTCCGTAGCACACAGTGCGAGCAGCAAAACAT 1020
L L P G S . S K . F A S . F A O R C C H C Y R H R G V T L V V V Y

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• TGGCTTCATTACAGCTCCGGTTCCCAACGATCAAGGCGAGTTACATGATCCCCATGTTGTGCAAAAAAGCGGTAGCTCCTTCGGTCTCCGATCGTTGT
ACCGAAGTAAGTCGAGGCCAAGGGTTGCTAGTTCGGCTCAATGTACTAGGGGGTACAACACGTTTTTTCGCCAATCGAGGAAGCCAGGAGGCTAGCAACA 1030
G F I Q L R F P T I K A S Y M I P H V V Q K S G . L L R S S D R C
CAGAAGTAAGTTGGCCGAGTGTATCACTCATGGTTATGGCAGCACTGCATAATCTCTTACTGTCATGCCATCCGTAAGATGCTTTTCTGTGACTGGT
GTCTTCATTCAACCGGCGTCACAATAGTGAGTACCAATACCGTCGTGACGTATTAAGAGAATGACAGTACGGTAGGCATTCTACGAAAAGACACTGACCA 1040
Q K . V G R S V I T H G Y G S T A . F S Y C H A I R K M L F C D V
GAGTACTCAACCAAGTCATTCTGAGAATAGTGATGCGGCGACCGAGTTGCTCTTGCCCGGCGTCAATACGGGATAATACCGCGCCACATAGCAGAAGTT
CTCATGAGTTGGTTCAGTAAGACTCTTATCACATACGCCGCTGGCTCAACGAGAACGGGCGCAGTTATGCCCTATTATGGCGCGGTGTATCGTCTTGAA 1050
V L N Q V I L R I V Y A A T E L L L P G V N T G . Y R A T . Q N F
TAAAAGTGCTCATCATTTGAAAAACGTTCTTCGGGGCGAAAACTCTCAAGGATCTTACCGCTGTTGAGATCCAGTTCGATGTAACCCACTCGTGACCCAA
ATTTTCACGAGTAGTAACCTTTTGCAAGAAGCCCCGCTTTTGAGAGTTCTTAGAATGGCGACAACCTCAGGTCAAGCTACATTGGGTGAGCACGTGGGTT 1060
K S A H H V K T F F G A K T L K D L T A V E I Q F D V T H S C T Q
CTGATCTTCAGCATCTTTTACTTTTACCAGCGTTTCTGGGTGAGCAAAAACAGGAAGGCAAAATGCCGCAAAAAAGGGAATAAGGGCGACACGGAAATGT
GACTAGAAGTCGTAGAAAAATGAAGTGGTCGCAAGACCCACTCGTTTTTGTCTTCCGTTTTACGGCGTTTTTCCCTTATTCCCGCTGTGCCTTTACA 1070
L I F S I F Y F H Q R F V V S K N R K A K C R K K G N K G D T E M
TGAATACTCATACTCTTCTTTTCAATATTATTGAAGCATTATCAGGGTTATTGTCTCATGAGCGGATACATATTTGAATGTATTTAGAAAAATAAAC 1080
ACTTATGAGTATGAGAAGGAAAAAGTTATAATAACTTCGTAATAGTCCCAATAACAGAGTACTCGCCTATGTATAAACTTACATAAACTCTTTTATTG
L N T H T L P F S I L L K H L S G L L S H E R I H I . M Y L E K . T
AAATAGGGGTTCCGCGCACATTTCCCGAAAAAGTGCCACCTGACGTC 10847
TTTATCCCAAGGCGCGTGTAAAGGGGCTTTTCACGGTGGACTGCAG
N R G S A H I S P K S A T . R

10 TCATCGGTATTACCATGGTGAATCGCGTTTGTGGCAGTACATCAATGGGCGGTGGATAGCGGTTTGGCTACGATGATG
TTTCCAAAGTCTCCACCACCATTCAGTCAATGGGAGTTGTTTGTGGCACAATAACGGGGATTTGACCTACGAGGGAA
CGTGAACAACCTCGGCCCAATTGACGCCAAATGGGCGGTAGGCGGTACGGGTGGGAGGTCTATATAAGCAGAGCTC
TGTGGCTAACTAGAGAACCACCTGCTTACTGGCTTATCGAAATTAATACGCTACATATAGGGAGACCCAAAGCTT
GGTACCGGACTCCGATAGCCACTAGTACGGCCCGCAGTGTGCTGGAATTCGCAAGTATCCATCCACTACGCTGGCGG
15 CCGCGCCCATGACGACGCTCAAAATGTAGAAATGTATACCAATCTACACGGGATGGGCCAATCGGCACCTTTGAAAG
GGCAGCTTTATCAAAATGCTATTAGGATATTTCCAATGATTTTCGGCACTATCGACTGGTTTCTCAGCTTTATTAAT
GTGATCGTTCCGATCAACGAATTTCTCGCCTGCATTACGAAACGTTTGGCAAAATACACCTGGAATGG
CTCGAAACGGTGTCTCGACTACCTCGAAAAATCTGGGTCTCGACTGTCTCGAAAGTACCCAAACCGAATCGCA
CGGAAACCTTGGGTGCGAGTTCTCGAGTGTCTTCTGCTCTCCACCTCAACGAGAAGCTTCGGCACTGAA
20 AAGATTCAGAAAGAAATTTGAGGCAATACCCCACTCATTGTGCGACCGGGTTTGAATTACCTCGCCACGT
GTCCGACCGTCAAGAACCGCTTCAGCAACTACCCCAATTCGAATTTCCAATTTCCACAAATGTCAACATCGAGCTTCAG
ACTCGACAGTCAAGAAATTCGAAATTTGATTCATCAAAGATTGGTATCAAGCGCAAGACGCTTGGACTTAAACCA
CGCTCATCATCAACCACCTTCATCAAAATAACAAATTCATTCCGCTCGCTCGAGCGCTTCGAGTTGGCAATAATAAT
25 GTTGGCTCGACGATATCCCATCTCGGAAGAGGTAGAATCATCATCAAGCTACAGCTGTGATTTGCAATCAAAAC
CGAGCTACCTCCCAACTCCAAAACTGTTATGACCAACAAACCCAGTAGTTGCTGTGCTACAACTACAAAAATC
GGAAGCTCAAAGTATGCGGCTCGCAAGCGCTGAGCAGCCCAAACTTGCTTCTGTGGAAGACTTTGGAGCAAA
ACAAGAGCGCGATAAACGCGGTGGTGGTGGTGGTGAATGCTGAAATTAAGTTATTACGATAGCAAAACCCCAT
CTTCTCATCGAATAGCCCAACCTACGGAAGGCGGGCGGGTGGCTCAACACCAAACTTGTGCGAAAACT
30 GCTGCCCCAGTGAAGAGTTGGCTGAAGCGCGGACCAAGTAAAGTGGGAAGTGGCAGCTGTATGTGCGAAGTTT
GTACGGCAAAAGTTTCTACCGTAAAGCGGACCGCCCAATCATATCTCAACAGACTCGAAACGATGCTCAAAGA
GCAGTGAAGAAGTCCGATACGCTGGAATCAACGACGCTGCCAACGTCATCATCGCGGAAGTTCCCT
AAGCATGCAATCCACATCTTCAAAGATTTCAAGCTCAGACGAAAAAGTCTCGTCTACAGACGATCTTACTCTTAA
CGCTCCCATCGTACAGTATCAGACAGCCGATAGCGGCAACACCGGTTTCTCAAATATTATCAACAGGCTG
TTGAGGAAAAACCAACACTGGCAGTGAAGGAGTGAAGACGACGCAAAAGATCCCACTCAGCTGTTCG
35 CCACGTGACACCCGACCAACATCGGATGTTTGTAGTCCAATTTGCGCAATGAAGAGTTGACAAATGACCCCGT
GATATCTGAAAAACGAGAACTGAAAGGCTCAATCAATGAGCATGCACGACGCGAGCTTCCACCGTTCGAC
CTCTAAATCAGTTGTTCACATTAATGACTTCAATCCGACAACCCCAACGTCAGTATGTTCTGTAACAAAGG
AAAAATCACATCGCTCGTCAAGTCTTGGATGAGCAGTCGTCGCGGTGGAAGCTCAATGCTGCGGATCGGCTTCAC
40 CGTCCGCTCAGGTGACTCCGCGCAGAAAACTTCTGGTAATCATTCGCTGGAGAGAAGTGGGAAGAATAAG
ACATCAGAAATCCAGCGGCTACACCTCTGACGCGCGGTGTTGCGATGTGCGCCAAAATGAGGAGAAAGCTGAAAG
AATACGATGACATGACTCTGCTGAGCAGCAACGCGCTATGCTGACAAGTTCGAAGACAGTCTCTCTGTGCTGT
GGAATATCCGATCAACACGAGCTCGACGACATATCCACGGACGATTGTGCGGAGTAGACATGGCAACGATCGCT
45 CTCACCAATAGCGACTATTTCCCACTTTGTTGCCATCCCACTCTCTCTCAAAAGCCCGAGTCCCCAGCT
GGTCTCCACATCAGTCTGATTCTCGATCTCGAGCAGAACAGGAGAATGTGTACAAAGCTTCTGTCGACGCGGA
ACGAGCCCAAGTGGCGCGCTGCCACCTCAACCTTCGGACAACATTCGCTAAGTATCCCGGGGATCTCATCTCTA
TTCTCCACACTTACGTGTACGTTGATAGGACACAATGCTATGCACTCAGACACTGACGACGACCTCTCTC
50 ACAAACCCCAAGCTATTCAGGCAATTTCAATCACTTGATCGTAAATGCCACCTTCAAGAGTTACATCTCAGGATCC
TGAAAAATCCCCCGCAGATTCTGCCAAAGTGTAGATGGGATCCCAACTATCAGTGGCTAGCAGCAGCATATG
GATCTCTCAATGAGAAGTAGCAATGCTATTCGGGACATGGCAGCTGACTTGGAGTTGTACAAGAACACTGTC
GACTCATATCAACCAAGAAACAGGAGAACTGGAGCATTTGTTGATCTTTTGAGCAAAAGCTTAGAAAACTCACT
CAACACATTGTGATGCTCAAGTTGAAGCCTGAAGAGGCAATACGATTACGGCAGGACATTTGCTATTGAGGGA
TATTAGCAATCATCTTGCATCCAACCTCAGCTCATGCTATCGTCAAGAGCGGTGGTGAAGCTTCTTCTCAACCTCTCT
55 GCTCGTTTGGCAAGAACAAGAGAGCTGGATCCGCTCTCTCACTCTCAAGTTCAACAAAGAGAAGACCAAGAA
TACGACGAAGACATATGCCATCAATTTCCGGGCTCAAGGAACCTTGTGACAACTTGTGATGTTGATTTGATTTGAAG
CAAGAGCTCAAAAGACGCGATGTGCACTTTACGAAGCATCTTGGCTGCAACTCAATCAAGTTGATGTTGATTTGAAG
TGTCTTGAGGGAGACAGTGAACAGTTGAAACCGGAGAACGAAGATTAATCGTAGGATATCTTGCCATGTCT
ACGGTCCAGCCACTCGTCTTCTTCCCGCGCTCAATTCAGTATCTACGACAGTATTCGCTCAACATCAATCGAATGTTGAT
60 GCGGTGAGCAGTACATCAGTATGTAATCTTCCGCGGCTCAATTCAGTATCTACGACGATGAGTATGATGATGCA
GTGGACATCGCTGGAGAAATCAGTTGATGCTGTTAAACCGGACGAAGAGATTAATCGTAGGATATCTTGCCATGTCT
AACCCTCAGTCACTGGAAGACATGATGTTCTATTCTAGGACTATTGAAGTCTACCTATCCAGATTTGAT
GTGGAGCATCAACTGGAAFCGATGCTCGTGATTCTTCTTGGTATCAAAATGGTGAATTCGACGCGCTATT
65 GGAAGTCCACAACCATGATAACCGAGCTCAACTGACATCTTACTCTCAACATCAATCGGAATGTTCAATG
CACGGTGGCCGACAGAGTCCGCTAGACAGCTTGGTCTTATGATGCTTCTTCAAGCAAAATGATTTCTCAACT
CGTCAAGTCAATTTGACAGAGAGCTGCTGGTGTAGCTGGAGCACTGGAATTTGAAAGAGCAAACTGGCGA
AGACCTTGGCTGCTTATGATCTATTGCAACAAATCAATCCGAAGATAGTATGTTGTAATCAGCATTCCTGAAAA
CAATAAAGAAAGATTTGCTTCAAGTGGAAACGACGCTGGAAAAAGATCTTGAGAAAGAAAGATTCGATCGATTAAT
70 GTTGCATTTGATGATGACAGTCAACCGATTCAAATCCCTGAGCTGCAATTCACCAATTCACAAATTCGAA
GTAATTCGAAATCGCTCGAAGGATTCATCTCAGTTCACCTCGGACGACGGGCGGTAGAGGATGAGTATCGTCT
AAGTGTACAGTCCGATCAGAGCTCTTCAAATCATTGACTTCTCCAATGCTCTCAGGCGGTCAAAATTTT
ATTGAGAAAAAGAAATTTCTGTTGATGTGACAGTTGGTCCAGAGCATGCTGAAGTGTCTTCAACTGTCTGAGTGA

FIG. 1 CONTINUED.

5 TCCCGTGAATGGTTCATTGCGATTGTGGAATGAGAACTTCATTCCATATTTGGAACGTGTTGCTAGAGATGGCAAA
AAAAACCTTCGGTTCGCTGCCTTCCTTCGAGGATCCACCGACATCGTCTCTAAAAAATGGCCGTGGTTCGATGG
TGAAAACCCGGAGAAATGTGCTCAAACGCTCTCAACTCCAAGACCTCGTCCCGTCACTGCGCAATTCATCCCGAC
AACACTTCAATCCCTCGAGTCTGTTGATCCAATTGCATGCTACCAAGCATCAGACCATCGACAACATTTGAACAG
AAGACTCTAATCTTCTCCTGCTCTCCCGCTTCCTTATCTTCTGATCCGGTACCTGATGATTCCTCCCATTTTCC
CCCTTTTCCCGCAATTTCCGAGAACCTCCTGTTCCCTTTGTTCTAGTCTCCCGGGTGGCGACGCCGGAAGCG
ATTTAAAAAGCTTTTCTTTCGGAACATTTCCCATTTGCTCATTAATAGTCAAAATGAATAACAGTGTATGTACTT
AAAAAAAAAAAAAAAAAAGTTCGAGGGGGGGCCCTATTCTATAGTGTACCTAAATGCTAGAGCTCGGTGATCAG
10 CCTCGACTGTGCTTCTAGTTGCCAGCCATCTGTTGTTGCCCTCCCGCGTGCCTTCTTACCTCGGAAAGGT
GCCACTCCCACTGTCTTCTTAATAAAATGAGGAAATTCATCGCATTTGCTGAGTAGGTGTCTATTCTATTCTG
GGGGGTGGGGTGGGGCAGGACAGCAAGGGGGAGGATTGGGAAGACAATAGCAGGCATGCTGGGGATGCGGT
GGGCTCTATGGCTTCTGAGGCGGAAAGAACAGCTGGGGCTCTAGGGGGTATCCCAACGCGCCCTGTAGCGG
CGCATTAAGCGCGGGGGTGTGGTGGTTACGCGCAGCGTACCGCTACACTTGCAGCGCCCTAGCGCCCGC
TCCTTTCCGCTTCTTCCCTTCTTCTCCTGCGCAGCTTCCCGGGCTTCCCGCTCAAGCTCTAAATCGGGGATCC
15 CTTTAGGGTTCCGATTAGTGTCTTACGGCACCTCGACCCCAAAAACTTGATTAGGGTGTGATGGTTACGTTT
GGGCCATCGCCCTGATGACGGTTTTTCCGCTTGTGAGTTCAGCTTCTTAATAGTGGACTCTTGT
CCAACTGGAACAACACTCAACCTATCTCGGTCTATTCTTTGATTATAAGGGATTGTTGGGATTTCGGCCTA
TTGGTTAAAAATGAGCTGATTTAAACAAAAATTAACGCGAATTAATTCTGTGGAATGTGTGTCAGTTAGGGTGT
GGAAAGTCCCGAGGCTCCCGAGGACAGGATGCAAAAGCATGCATCTCAATTAGTCAGCAACCAAGGTGT
20 GGAAAGTCCCGAGGCTCCCGAGGACAGGATGCAAAAGCATGCATCTCAATTAGTCAGCAACCAATGTCC
GCCCTAATCTCCGCCATCCCGCCCTAATCTCCGCCAGTTCGCCCATCTCTCCGCCCATGGCTGACTAATTT
TTTTATTTATGACAGGGCGAGGCGCCCTCTGCTCTGAGCTATTCCAGAAAGTAGTGAGGAGGCTTTTGGGA
GGCCTAGGCTTTTGCAAAAAGCTCCCGGGAGCTTGATATCCATTTCCGATCTGATCAAGAGACAGGATGAGG
ATCGTTTCGCTGATTTGAACAAGATGGATTGCACGCGAGTTCTCCGGCCGCTTGGGTGGAGAGGCTATTCCGGC
25 TATGACTGGGCACAACAGACAATCGGCTGCTGTATGCCCGCGTGTTCGGCTGTACAGCGAGGGCGCCCG
GTCTTTTTGTCAAGACCGACCTGTCCGGTGCCTGAATGAACTGCAGGACGAGGACGCGGCTATCGTGGC
TGGCCACGACGGGGCTTCTTGGCGAGCTGTGCTCGACGTTGTCACTGAAGCGGGAAGGGAGTGGCTGCTATT
GGCGGAAGTGGCGGGGAGGATCTCCTGTCTACCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT
30 GCAATGCGCGGGCTGCATACGCTTGATCCGGCTACCTGCCATTTCGACCACCAAGCGAAACATCGCATCGAGC
GAGCAGTACTCGGATGGAAGCCGGCTTGTGATCAGGATGATCTGGACGAAGAGCATCAGGGGCTCGGC
CAGCCGAATCTGTCGCCAGGCTCAAGGGCGCATGCCCCGACGGGAGGATCTCGTGCATGCCATGGCGGAT
CCTGCTTGGCGAATATCATGTTGGAAATGGCCGCTTTCTGGATTCTCGACTGTGGCCGGCTGGGTGTGGC
GGACCGCTACAGGACATAGCGTTGGCTACCCGATGATTTGCTGAAGAGCTTGGCGGGCAATGGGCTGACCGC
35 TTCTCGTGTCTTACGGTATCGCCGCTCCCGATTCCGACGCGCATCGCTTCTATCGCTTCTTACGAGTCTT
CTGAGCGGACTCTGGGGTTCGAAATGACCGACCAAGCGACGCCCAACCTGCCATCAGGAGATTTCGATTCCA
CCCGCGCTTCTATGAAAGTTGGGCTTCGGAATCGTTTTCCGGGACGCCGCTGGATCTCCCTACCGCGCG
GGATCTCATGCTGGAGTTCTTCCGCCACCCCAACTTGTATTGACGCTTATAATGGTTACAAATAAGCAATAG
CATCACAAAATTCACAAATAAGCAATTTTTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT
40 ATCATGCTGTATACCGTGCACCTCTAGCTAGAGCTTGGCGTAATCATGCTCATAGCTGTTCTGTGTGAAAT
GTTATCCGCTCACAAATCCACACAACATACGAGCGGGAAGCATAAAGTGTAAAGCTTGGGGTGCCTAATGAGTG
AGCTAATCATCAATTAATGCTTGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT
ATGAATCGGCCCAACGCGCGGGGAGAGGCGGTTTGGCTATTGGGCGCTCTTCCGCTTCTCGCTCACTGCTCG
45 GCTGCGGTTTTCCATAGGCTCCGCCCGCTGACGAGCATCAAAAAATCGACGCTCAAGTCAGAGGTGGCGCA
AACCAGCAGGACTATAAAGATACAGGCGTTTCCCGCTGGAAGCTCCTCGTGGCTCTCTGTTCCGACCT
GCCGCTTACCGGATACCTGTCCGCTTCTCCTTCCGGGAAGCGTGGCGCTTCTCAATGCTCAGCGTGTAGG
TATCTCAGTTCCGTTGAGGTGCTTCCGCTCAAGCTGGGCTGTGTGACGAACCCCCGTTACGCCCCGACCGCT
50 CGGCTTATCCGGTAACTATCGTCTTGAAGTCAACCCGGAAGACAGGCTTATCGCCACTGGCAGGCACT
GGTAACAGGATTAGCAGAGCGAGGATGTAGGCGGTGCTACAGAGTTCTTGAAGTGGTGGCCTAACTACGGCT
ACACTAGAAGGACAGTATTGGTATCTGCGCTCTGCTGAAGCCAGTTACCTTCGGAAGAGGTTGGTAGCTCT
TGATCCGGCAACAAACACCGCTGGTAGCGGTGGTTTTTTGTTTGAAGCAGCAGATTACGGCGAGAAAAA
AGGATCTCAAGAGATCTTTGATCTTTCTACGGGGTCTGACGCTCAGTGGAAACGAAACTCAGCTTAAGGGA
55 TTTTGGTCATGAGATTATCAAAAAGGATCTTACCTAGATCTTTTAAATTAATAATGAAGTTTAAATCAATCTAA
AGTATATAGTAACTTGGTCTGACAGCTTACCAATGCTTAACTAGTGAGGACCTATCTCAGCGATCTGTCTA
TTTCTGTTATCATAGTTGCTGACTCCCGCTGCTGTAGATACTACGATACGGGAGGGCTTACCATCTGCGCC
CAGTGTGCAATGATACCGCGAGACCCAGCTACCGGCTCCAGATTTATCAGCAATAAACACGCGACCGGAA
GGCGCGAGCGCAGAAAGTGGTCTGCACTTTATCCGCTCCATCCAGTCTAATTTGTTGCCGGGAAGGTAGTA
60 GTAAGTAGTTCGCCAGTTAATAGTTTGGCAACGTTTGGCATTTGCTACAGGATCGTGGTTCAGCGCTGCT
GTTTGGTATGGCTTCTTACGCTCCGCTTCCCAACGATCAAGGCGAGTTACATGATCCCCCATGTTGTGCAAAA
AAGCGGTAGCTCTTCCGCTCCGATGCTGTCAGAAATAGTTTGGCGCAGTGTATCTACTCATGTTATG
GCAGCACTGCATAATCTCTTACTGTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT
65 TCATTCTGAGAAATAGTATGCGGGCAGCGAGTTGCTTTCGCCGGCGTCAATACGGGATATACCGCGCCACA
TAGCAGAACTTTAAAGTGCTCATCTATTGGAACGTTCTTCCGGGGCAAAACTCTCAAGGATATCTGCTCCGCT
GAGATCCAGTTGATGTAACCCACTGCTGCACCCAACTGATCTTACGATCTTTTACTTTACCGGGTTCTGCG
GTGAGCAAAAACAGGAAGGCAAAATGCCGCAAAAAGGGAATAAGGGCGACAGGAAATGTTGAATCTATAC
TCTTCTTTTCAATATTTATGAAGCATTTATCAGGGTTATTGTCTCATGAGCGGATACATTTGAATGATTAG
AAAAATAACAAATAGGGGTTCCGGCGCACATTTCCCGGAAAAGTGCCACCTGACGTCGACGGATCGGGGATCT
70 CCGGATCCCTATGGTGCATCTCAGTACAATCTGCTCTGATGCGCGCATAGTTAAGCCAGTATCTGCTCCGCT
TTGTGTGTTGGAGGTGCTGAGTAGTGGCGAGCAAAATTAAGCTACAAAGGCAAGGCTTACCGGCAAT
GCATGAAGAATCTGCTTAGGGTTAGGCGTTTGGCTGCTTCCGATGACGGGCGAGATACGCGTTGACAT
TGATTATTGACTAGTTATTAATAGTAATCAATTACGGGGTCAATAGTTATAGCCCATATATGAGGTTCCGCTTA
75 CATAACTTACGGTAAATGGCCCGCTGGCTGACCGCCCAACGACCCCGCCCATGACGTCATTAATGACGTAT
GTTCCCATAGTAACGGCAATAGGAGCTTCCATTGACGTCATAGGGTGGACTATTTACGGTAACTGCCACCTT
GGCAGTACATCAAGTGTATCATATGCCAAGTACGCCCCCTATTGACGTCATGACGGTA

FIG. 2.

Met Thr Thr Ser Asn Val Glu Leu Ile Pro Ile
Tyr Thr Asp Trp 15

Ala Asn Arg His Leu Ser Lys Gly Ser Leu Ser
Lys Ser Ile Arg 30

Asp Ile Ser Asn Asp Phe Arg Asp Tyr Arg Leu
Val Ser Gln Leu 45

Ile Asn Val Ile Val Pro Ile Asn Glu Phe Ser
Pro Ala Phe Thr 60

Lys Arg Leu Ala Lys Ile Thr Ser Asn Leu Asp
Gly Leu Glu Thr 75

Cys Leu Asp Tyr Leu Lys Asn Leu Gly Leu Asp
Cys Ser Lys Leu 90

Thr Lys Thr Asp Ile Asp Ser Gly Asn Leu Gly
Ala Val Leu Gln 105

Leu Leu Phe Leu Leu Ser Thr Tyr Lys Gln Lys
Leu Arg Gln Leu 120

Lys Lys Asp Gln Lys Lys Leu Glu Gln Leu Pro
Thr Ser Ile Met 135

Pro Pro Ala Val Ser Lys Leu Pro Ser Pro Arg
Val Ala Thr Ser 150

Ala Thr Ala Ser Ala Thr Asn Pro Asn Ser Asn
Phe Pro Gln Met 165

Ser Thr Ser Arg Leu Gln Thr Pro Gln Ser Arg
Ile Ser Lys Ile 180

Asp Ser Ser Lys Ile Gly Ile Lys Pro Lys Thr
Ser Gly Leu Lys 195

Pro Pro Ser Ser Ser Thr Thr Ser Ser Asn Asn
Thr Asn Ser Phe 210

Arg Pro Ser Ser Arg Ser Ser Gly Asn Asn Asn
Val Gly Ser Thr 225

Ile Ser Thr Ser Ala Lys Ser Leu Glu Ser Ser
Ser Thr Tyr Ser 240

FIG. 2 CONTINUED.

Ser Ile Ser Asn Leu Asn Arg Pro Thr Ser Gln
Leu Gln Lys Pro 255

Ser Arg Pro Gln Thr Gln Leu Val Arg Val Ala
Thr Thr Thr Lys 270

Ile Gly Ser Ser Lys Leu Ala Ala Pro Lys Ala
Val Ser Thr Pro 285

Lys Leu Ala Ser Val Lys Thr Ile Gly Ala Lys
Gln Glu Pro Asp 300

Asn Ser Gly Gly Gly Gly Gly Gly Met Leu Lys
Leu Lys Leu Phe 315

Ser Ser Lys Asn Pro Ser Ser Ser Ser Asn Ser
Pro Gln Pro Thr 330

Arg Lys Ala Ala Ala Val Pro Gln Gln Gln Thr
Leu Ser Lys Ile 345

Ala Ala Pro Val Lys Ser Gly Leu Lys Pro Pro
Thr Ser Lys Leu 360

Gly Ser Ala Thr Ser Met Ser Lys Leu Cys Thr
Pro Lys Val Ser 375

Tyr Arg Lys Thr Asp Ala Pro Ile Ile Ser Gln
Gln Asp Ser Lys 390

Arg Cys Ser Lys Ser Ser Glu Glu Glu Ser Gly
Tyr Ala Gly Phe 405

Asn Ser Thr Ser Pro Thr Ser Ser Ser Thr Glu
Gly Ser Leu Ser 420

Met His Ser Thr Ser Ser Lys Ser Ser Thr Ser
Asp Glu Lys Ser 435

Pro Ser Ser Asp Asp Leu Thr Leu Asn Ala Ser
Ile Val Thr Ala 450

Ile Arg Gln Pro Ile Ala Ala Thr Pro Val Ser
Pro Asn Ile Ile 465

Asn Lys Pro Val Glu Glu Lys Pro Thr Leu Ala
Val Lys Gly Val 480

Lys Ser Thr Ala Lys Lys Asp Pro Pro Pro Ala
Val Pro Pro Arg 495

Asp Thr Gln Pro Thr Ile Gly Val Val Ser Pro
Ile Met Ala His 510

FIG. 2 CONTINUED.

Lys Lys Leu Thr Asn Asp Pro Val Ile Ser Glu
Lys Pro Glu Pro 525

Glu Lys Leu Gln Ser Met Ser Ile Asp Thr Thr
Asp Val Pro Pro 540

Leu Pro Pro Leu Lys Ser Val Val Pro Leu Lys
Met Thr Ser Ile 555

Arg Gln Pro Pro Thr Tyr Asp Val Leu Leu Lys
Gln Gly Lys Ile 570

Thr Ser Pro Val Lys Ser Phe Gly Tyr Glu Gln
Ser Ser Ala Ser 585

Glu Asp Ser Ile Val Ala His Ala Ser Ala Gln
Val Thr Pro Pro 600

Thr Lys Thr Ser Gly Asn His Ser Leu Glu Arg
Arg Met Gly Lys 615

Asn Lys Thr Ser Glu Ser Ser Gly Tyr Thr Ser
Asp Ala Gly Val 630

Ala Met Cys Ala Lys Met Arg Glu Lys Leu Lys
Glu Tyr Asp Asp 645

Met Thr Arg Arg Ala Gln Asn Gly Tyr Pro Asp
Asn Phe Glu Asp 660

Ser Ser Ser Leu Ser Ser Gly Ile Ser Asp Asn
Asn Glu Leu Asp 675

Asp Ile Ser Thr Asp Asp Leu Ser Gly Val Asp
Met Ala Thr Val 690

Ala Ser Lys His Ser Asp Tyr Ser His Phe Val
Arg His Pro Thr 705

Ser Ser Ser Ser Lys Pro Arg Val Pro Ser Arg
Ser Ser Thr Ser 720

Val Asp Ser Arg Ser Arg Ala Glu Gln Glu Asn
Val Tyr Lys Leu 735

Leu Ser Gln Cys Arg Thr Ser Gln Arg Gly Ala
Ala Ala Thr Ser 750

Thr Phe Gly Gln His Ser Leu Arg Ser Pro Gly
Tyr Ser Ser Tyr 765

Ser Pro His Leu Ser Val Ser Ala Asp Lys Asp
Thr Met Ser Met 780

FIG. 2 CONTINUED.

His Ser Gln Thr Ser Arg Arg Pro Ser Ser Gln
Lys Pro Ser Tyr 795

Ser Gly Gln Phe His Ser Leu Asp Arg Lys Cys
His Leu Gln Glu 810

Phe Thr Ser Thr Glu His Arg Met Ala Ala Leu
Leu Ser Pro Arg 825

Arg Val Pro Asn Ser Met Ser Lys Tyr Asp Ser
Ser Gly Ser Tyr 840

Ser Ala Arg Ser Arg Gly Gly Ser Ser Thr Gly
Ile Tyr Gly Glu 855

Thr Phe Gln Leu His Arg Leu Ser Asp Glu Lys
Ser Pro Ala His 870

Ser Ala Lys Ser Glu Met Gly Ser Gln Leu Ser
Leu Ala Ser Thr 885

Thr Ala Tyr Gly Ser Leu Asn Glu Lys Tyr Glu
His Ala Ile Arg 900

Asp Met Ala Arg Asp Leu Glu Cys Tyr Lys Asn
Thr Val Asp Ser 915

Leu Thr Lys Lys Gln Glu Asn Tyr Gly Ala Leu
Phe Asp Leu Phe 930

Glu Gln Lys Leu Arg Lys Leu Thr Gln His Ile
Asp Arg Ser Asn 945

Leu Lys Pro Glu Glu Ala Ile Arg Phe Arg Gln
Asp Ile Ala His 960

Leu Arg Asp Ile Ser Asn His Leu Ala Ser Asn
Ser Ala His Ala 975

Asn Glu Gly Ala Gly Glu Leu Leu Arg Gln Pro
Ser Leu Glu Ser 990

Val Ala Ser His Arg Ser Ser Met Ser Ser Ser
Ser Lys Ser Ser 1005

Lys Gln Glu Lys Ile Ser Leu Ser Ser Phe Gly
Lys Asn Lys Lys 1020

Ser Trp Ile Arg Ser Ser Leu Ser Lys Phe Thr
Lys Lys Lys Asn 1035

Lys Asn Tyr Asp Glu Ala His Met Pro Ser Ile
Ser Gly Ser Gln 1050

FIG. 2 CONTINUED.

Gly Thr Leu Asp Asn Ile Asp Val Ile Glu Leu
Lys Gln Glu Leu 1065

Lys Glu Arg Asp Ser Ala Leu Tyr Glu Val Arg
Leu Asp Asn Leu 1080

Asp Arg Ala Arg Glu Val Asp Val Leu Arg Glu
Thr Val Asn Lys 1095

Leu Lys Thr Glu Asn Lys Gln Leu Lys Lys Glu
Val Asp Lys Leu 1110

Thr Asn Gly Pro Ala Thr Arg Ala Ser Ser Arg
Ala Ser Ile Pro 1125

Val Ile Tyr Asp Asp Glu His Val Tyr Asp Ala
Ala Cys Ser Ser 1140

Thr Ser Ala Ser Gln Ser Ser Lys Arg Ser Ser
Gly Cys Asn Ser 1155

Ile Lys Val Thr Val Asn Val Asp Ile Ala Gly
Glu Ile Ser Ser 1170

Ile Val Asn Pro Asp Lys Glu Ile Ile Val Gly
Tyr Leu Ala Met 1185

Ser Thr Ser Gln Ser Cys Trp Lys Asp Ile Asp
Val Ser Ile Leu 1200

Gly Leu Phe Glu Val Tyr Leu Ser Arg Ile Asp
Val Glu His Gln 1215

Leu Gly Ile Asp Ala Arg Asp Ser Ile Leu Gly
Tyr Gln Ile Gly 1230

Glu Leu Arg Arg Val Ile Gly Asp Ser Thr Thr
Met Ile Thr Ser 1245

His Pro Thr Asp Ile Leu Thr Ser Ser Thr Thr
Ile Arg Met Phe 1260

Met His Gly Ala Ala Gln Ser Arg Val Asp Ser
Leu Val Leu Asp 1275

Met Leu Leu Pro Lys Gln Met Ile Leu Gln Leu
Val Lys Ser Ile 1290

Leu Thr Glu Arg Arg Leu Val Leu Ala Gly Ala
Thr Gly Ile Gly 1305

Lys Ser Lys Leu Ala Lys Thr Leu Ala Ala Tyr
Val Ser Ile Arg 1320

FIG. 2 CONTINUED.

Thr Asn Gln Ser Glu Asp Ser Ile Val Asn Ile
Ser Ile Pro Glu 1335

Asn Asn Lys Glu Glu Leu Leu Gln Val Glu Arg
Arg Leu Glu Lys 1350

Ile Leu Arg Ser Lys Glu Ser Cys Ile Val Ile
Leu Asp Asn Ile 1365

Pro Lys Asn Arg Ile Ala Phe Val Val Ser Val
Phe Ala Asn Val 1380

Pro Leu Gln Asn Asn Glu Gly Pro Phe Val Val
Cys Thr Val Asn 1395

Arg Tyr Gln Ile Pro Glu Leu Gln Ile His His
Asn Phe Lys Met 1410

Ser Val Met Ser Asn Arg Leu Glu Gly Phe Ile
Leu Arg Tyr Leu 1425

Arg Arg Arg Ala Val Glu Asp Glu Tyr Arg Leu
Thr Val Gln Met 1440

Pro Ser Glu Leu Phe Lys Ile Ile Asp Phe Phe
Pro Ile Ala Leu 1455

Gln Ala Val Asn Asn Phe Ile Glu Lys Thr Asn
Ser Val Asp Val 1470

Thr Val Gly Pro Arg Ala Cys Leu Asn Cys Pro
Leu Thr Val Asp 1485

Gly Ser Arg Glu Trp Phe Ile Arg Leu Trp Asn
Glu Asn Phe Ile 1500

Pro Tyr Leu Glu Arg Val Ala Arg Asp Gly Lys
Lys Thr Phe Gly 1515

Arg Cys Thr Ser Phe Glu Asp Pro Thr Asp Ile
Val Ser Lys Lys 1530

Trp Pro Trp Phe Asp Gly Glu Asn Pro Glu Asn
Val Leu Lys Arg 1545

Leu Gln Leu Gln Asp Leu Val Pro Ser Pro Ala
Asn Ser Ser Arg 1560

Gln His Phe Asn Pro Leu Glu Ser Leu Ile Gln
Leu His Ala Thr 1575

Lys His Gln Thr Ile Asp Asn Ile

FIG. 3. : tblastn search of the EST division of Genbank with the ORF of the longest known *Ce*-UNC-53 cDNA, tb3-M5, reveals two EST's with homology to a predicted coiled-coil region in *Ce*-UNC-53.

TBLASTN 1.4.9MP [26-March-1996] [Build 14:27:13 Apr 1 1996]

Reference: Altschul, Stephen F., Warren Gish, Webb Miller, Eugene W. Myers, and David J. Lipman (1990). Basic local alignment search tool. J. Mol. Biol. 215:403-10.

Query= tb3 M5 ORF

(1583 letters)

Database: Non-redundant Database of GenBank EST Division
647,253 sequences; 234,216,808 total letters.

| Sequences producing High-scoring Segment Pairs: | Reading Frame | High Score | Smallest Sum P(N) | Probability N |
|-----------------------------------------------------------|---------------|------------|-------------------|---------------|
| dbj D35780 CELK025D6F C.elegans cDNA clone yk25d6 : 5'... | +2 | 358 | 7.9e-54 | 3 |
| dbj D33048 CELK025D6R C.elegans cDNA clone yk25d6 : 3'... | -1 | 177 | 8.6e-16 | 1 |
| gb H09036 H09036 y196c11.r1 Homo sapiens cDNA clo... | +1 | 115 | 1.1e-05 | 1 |
| gb AA049124 AA049124 mj46f04.r1 Soares mouse embryo N... | +3 | 106 | 8.6e-05 | 1 |
| gb R91475 R91475 yq08c11.r1 Homo sapiens cDNA clo... | +2 | 59 | 0.21 | 2 |
| gb T23446 T23446 seq2955 Homo sapiens cDNA clone ... | -1 | 61 | 0.99 | 2 |
| gb R86390 R86390 SW13CA339SK Brugia malayi infect... | +2 | 74 | 0.996 | 1 |
| gb T44781 T44781 8044 Arabidopsis thaliana cDNA c... | +1 | 71 | 0.9992 | 1 |
| gb T75582 T75582 yd63f11.r1 Homo sapiens cDNA clo... | +2 | 64 | 0.99992 | 2 |

30 gb|H09036|H09036 y196c11.r1 Homo sapiens cDNA clone 46037 5'.
Length = 489

Plus Strand HSPs:

Score = 115 (52.1 bits), Expect = 1.1e-05, P = 1.1e-05

35 Identities = 22/70 (31%), Positives = 45/70 (64%), Frame = +1

Query: 1059 IELKQELKERDSALYEVRLDNLDRAREVDVLRETIVNKLKTENKOLKKEVDKLTNGPATRA 1118
++L+ EL+... L ++RL+ L A ++D LRE +N++++E ++LK D+L +

Sbjct: 7 HOLRNLRLDKEMKLTDIRLEALSSAHOLDOLREAHNRMQSEIEKLKXNDRLKSESOQSG 186

40 Query: 1119 SSRASIPVIY 1128

SR S P ++

Sbjct: 187 CSRGSPSVH 216

45 gb|AA049124|AA049124 mj46f04.r1 Soares mouse embryo NbME13.5 14.5 Mus
musculus cDNA clone 479167 5'.
Length = 337

Plus Strand HSPs:

Score = 106 (48.0 bits), Expect = 8.6e-05, P = 8.6e-05

50 Identities = 23/58 (39%), Positives = 38/58 (65%), Frame = +3

Query: 1057 DVIELKQELKERDSALYEVRLDNLDRAREVDVLRETIVNKLKTENKOLKKEVDKLTNGP 1114
+V EL+ EL E++ L ++RL+ L A ++D LAET++ E LK E D+L P

55 Sbjct: 99 EVSELASELWEKEMKLTDIRLEALNSAHOLDOLRETVMHNLQEVOLLKAENDRLKVP 272

FIG. 4.

A Search of the Genbank databases with part of the nucleotide binding domain of *Ce-UNC-53* does not identify statistically significant proteins except for the *C. elegans* cosmid containing *Ce-unc-53*.

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5  TBLASTN 1.4.8MP [20-June-1995] [Build 18:00:05 Aug 29 1995]
   Query= section5 (240 letters)
   >lcl|section5
   ILTERRLVLAGATGIGKSKLAKTLAAYVSIRTNOSEDSIVNISIPENNKELLQVERRLE
10  KILRSKESCIVILDNIPKNRIAFVVSFANVPLONNEGFVVCTVNRYOIPELOIHNNFK
   MSVMSNRLEGFILRYLRRRAVEDEYRLTVQMPSELFKKIIDFFPIALQAVNNFIKTNVSVO
   VTVGPRACLNCLTVGGSREWFIRLWNNENFIPYLERVAROGKKNLRLSLHFLRGSHRHRLX

   Database: Non-redundant PDB+GBupdate+GenBank+EMBLupdate+EMBL
             520,383 sequences: 367,017,413 total letters.

15                                     Smallest
                                     Sum
                                     Probability
Sequences producing High-scoring Segment Pairs:
emb1247810|CEFA45E10 Caenorhabditis elegans cosmid F45... -2 1131 5.1e-158 2
gb1R41071|R41071 Hk575-f Homo sapiens cDNA clone k... +2 53 0.33 2
20  gb1T44781|T44781 8044 Arabidopsis thaliana cDNA cl... +1 74 0.35 1
   emb1248334|CEP10B5 Caenorhabditis elegans cosmid F10... +3 71 0.83 3
   gb1M81884|EPFPCPG Epifagus virginiana chloroplast c... +1 49 0.91 4
   gb1L09547|PEAPCLP Pisum sativum (clone pCLp) nuclea... +1 71 0.99 1
25  gb1M32604|TOMCD4B Tomato ATP-dependent protease (CD... +1 71 0.99 1
   emb1X69188|APIUSGA A.pyhilitidis mRNA for gamma-tubulin +2 56 0.992 3
   gb1T44782|T44782 8044 Arabidopsis thaliana cDNA cl... +1 68 0.9995 1
   gb1M17087|HUMRASK12 Human c-ras-K1-2 activated oncoge... +2 58 0.9998 1
   emb1X57702|IGGNATRIUP G.gallus RNA for precursor of nat... +3 56 0.9999 2
30  gb1K01520|HUMRASKB1 Human lung adenocarcinoma (PR371)... +2 57 0.99995 1

>gb1R41071|R41071 Hk575-f Homo sapiens cDNA clone k575-f.
   Length = 310
   Plus Strand MSPs:

35  Score = 53 (24.5 bits), Expect = 0.40, Sum P(2) = 0.33
   Identities = 9/15 (60%), Positives = 13/15 (86%), Frame = +2

Query: 130 GFILRYLRRRAVEDE 144
      GF+RYLRR+ VE +
40  Sbjct: 26 GFLVRYLRRLVESD 70

   Score = 47 (21.7 bits), Expect = 0.40, Sum P(2) = 0.33
   Identities = 9/26 (34%), Positives = 17/26 (65%), Frame = +3

45  Query: 170 NNFIEKTNVSVDVTVGPRACLNCLTV 195
      F-EK ---O -GP L+ PL +
   Sbjct: 147 HTFLEKXSTLDFLIGPCFFLSGPLAL 224

```

FIG. 5.

Three frame translation of EST gb:R41071.

Regions of homology region with *Ce-Unc-53* in two different frames are underlined. The spacing between the blocks of homology is of similar size as that in *Ce-UNC-53*.

- 5 Subsequent re-cloning and re-sequencing of this region in man identified multiple sequencing errors gb:R41071, and identified an ORF which is more homologous to and co-linear with *Ce-UNC-53* (see alignment in fig 12).

```

10
CTCCAACAACGTGGAGCCAGCCAATGGCTTCCTGGTTCGTTACCTGAGGAGGAAGCTGGT
    10      20      30      40      50      60
15  L Q Q R G A S Q W L P G S L P E E A G
   S N N V E P A N G F L V R Y L R R K L V
   P T T W S Q P M A S W F V T * G G S W *

AGAGTCAGACAGCGACATCAATGCCAACAAGGAAGAGCTGCTTCGGGGTGCTCGACTTGG
    70      80      90      100     110     120
20  R V R Q R H Q C Q Q G R A A S G C S T W
   E S D S D I N A N K E E L L R G A R L G
   S Q T A T S M P T R K S C F G V L D L G

GTACCCAAGCCTGTGGTATCATCTTCCACACCTTCCTTGAGAAGCACAGCACCTTAGACT
    130     140     150     160     170     180
25  V P K P V V S S S T P S L R S T A P * T
   Y P S L W Y H L P H L P * E A Q H L R L
   T Q A C G I I F H T F L E K H S T L D F

30  TTCTCATCGGCCCTTGCTTCTTTCTGTGCGGTCCATTGGCATTGAGGCTTCCGGACCTTG
    190     200     210     220     230     240
   F S S A L A S F C R V H W H * G F R T L
   S H R P L L L S V G S I G I E A S G P C
   L I G P C F F L S G P L A L R L P D L V

35  TTTATTGACCTGTGGACAACCTATCATTTCTATCTACAGGAGGAGCCAAGGATTGGAT
    250     260     270     280     290     300
   F I D L W T T L S F P I Y R R S Q G L D
   L L T C G Q L Y H F L S T G G A K D W I
   Y * P V D N S I I S Y L Q E E P R I G *

40  AAAGGTCCAT
    310
   K G P
   K V H
45  R S

```

FIG. 6 : blastn search of the EST division of Genbank with Hu-unc-53/1 cDNA 3b.

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BLASTN 1.4.9MP (26-March-1996) (Build 14:27:07 Apr 1 1996)
Query= Hu-unc-53/1 cDNA 3b
      (3256 letters)
Database: Non-redundant Database of GenBank EST Division
          647,253 sequences; 234,216,808 total letters.

```

| Sequences producing High-scoring Segment Pairs: | High Score | Smallest Sum | Probability | LOCUS |
|--------------------------------------------------------------|------------|--------------|--------------|-------------|
| | | P(N) | N assignment | |
| gb N36659 N36659 yx91b09.r1 Homo sapiens cDNA clone 2... | 1668 | 2.1e-130 | 1 | hu-UNC- |
| 53/1 | | | | |
| gb AA043997 AA043997 zk58a01.r1 Soares pregnant uterus Nb... | 1316 | 8.3e-129 | 3 | hu-UNC-53/1 |
| gb AA049124 AA049124 mj46204.r1 Soares mouse embryo NbME1... | 1324 | 9.1e-102 | 1 | ms-UNC-53/1 |
| gb T05560 T05560 ESTC3449 Homo sapiens cDNA clone HFB... | 892 | 5.1e-84 | 3 | hu-UNC-53/1 |
| gb N24681 N24681 yx91b09.s1 Homo sapiens cDNA clone 2... | 782 | 9.9e-75 | 2 | hu-UNC-53/1 |
| gb R41071 R41071 Hk575-f Homo sapiens cDNA clone k575-f... | 535 | 1.5e-72 | 4 | hu-UNC-53/1 |
| gb N89104 N89104 K7846F Fetal heart, Lambda ZAP Expre... | 451 | 7.3e-57 | 2 | hu-UNC-53/1 |
| gb R41073 R41073 Hk144-f Homo sapiens cDNA clone k144-f... | 555 | 1.5e-36 | 1 | hu-UNC-53/1 |
| gb R15492 R15492 HH434-F Homo sapiens cDNA clone H434-F... | 416 | 2.3e-29 | 2 | hu-UNC-53/1 |
| gb H09036 H09036 y196c11.r1 Homo sapiens cDNA clone 4... | 438 | 9.4e-26 | 1 | hu-UNC-53/2 |
| gb W91567 W91567 MTA.C36.093.A MTA adult mouse thymus... | 317 | 1.9e-17 | 2 | ms-UNC-53/7 |
| gb W74400 W74400 zd62d10.r1 Soares fetal heart NbHH19... | 243 | 2.2e-09 | 1 | hu-UNC-53/1 |
| gb AA003314 AA003314 mg56h10.r1 Soares mouse embryo NbME1... | 141 | 0.54 | 1 | |

FIG. 7.

TBLASTN search of the Genbank sequence database with the 961 aminoacid ORF of cDNA 3b of hu-UNC-53/1. hu-UNC-53/1 forms a unique pair with *Ce*-UNC-53 (cosmid F45E10) compared to the rest of the database.

5

10

TBLASTN 1.4.3MP (26-March-1996) (Build 14:27:13 Apr 1 1996)

15 Query= tmpseq_1
(961 letters)

Database: Non-redundant GenBank+EMBL+DDBJ+PDB sequences
261,674 sequences; 371,416,172 total letters.

20

| Sequences producing High-scoring Segment Pairs: | Reading Frame | High Score | Smallest Sum Probability P(N) | N |
|--------------------------------------------------------------------|------------------|---------------|----------------------------------------|---|
| 25 emb1247810 CEF45E10 <i>Caenorhabditis elegans</i> cosmid F45E10 | -2 | 158 | 2.3e-32 | 7 |
| gb1M97501 HUMCLIP Human cytoplasmic linker protein-... | +3 | 83 | 0.47 | 1 |
| emb1X64838 HSRESTIN <i>H. sapiens</i> mRNA for restin | +1 | 83 | 0.47 | 1 |
| gb1M58752 ECOMCRBC <i>E. coli</i> mcrB and mcrC genes, compl... | +3 | 82 | 0.56 | 1 |
| emb1211582 SCNUF1G <i>S. cerevisiae</i> nuf1 gene | -1 | 82 | 0.61 | 2 |
| 30 emb1X73297 SCSETRP4 <i>S. cerevisiae</i> spacer element | -1 | 82 | 0.74 | 2 |
| emb1X54002 XLKINESIN <i>X. laevis</i> mRNA for kinesine | +2 | 63 | 0.85 | 5 |
| gb1U42409 DDU42409 <i>Dictyostelium discoideum</i> myosin h... | +3 | 66 | 0.92 | 2 |
| gb1U10399 YSC8082 <i>Saccharomyces cerevisiae</i> chromoso... | +2 | 77 | 0.93 | 2 |
| 35 gb1U20810 ATU20810 <i>Arabidopsis thaliana</i> cytoskeleton... | +1 | 77 | 0.95 | 2 |
| gb1L07879 LEIKINLIKE <i>Leishmania chagasi</i> kinesin-like p... | +2 | 78 | 0.95 | 1 |
| gb1L03188 YSCINTANA <i>Saccharomyces cerevisiae</i> integrin... | +2 | 65 | 0.997 | 1 |
| gb1U28372 YSCD9476 <i>Saccharomyces cerevisiae</i> chromoso... | +3 | 82 | 0.9991 | 2 |
| gb1M94362 HUMLAMBA Human lamin B2 (LAMB2) mRNA, part... | +1 | 75 | 0.9996 | 1 |
| 40 gb1M58337 VACHAGMA <i>Vaccinia virus</i> hemagglutinin gene. | +1 | 74 | 0.99995 | 1 |

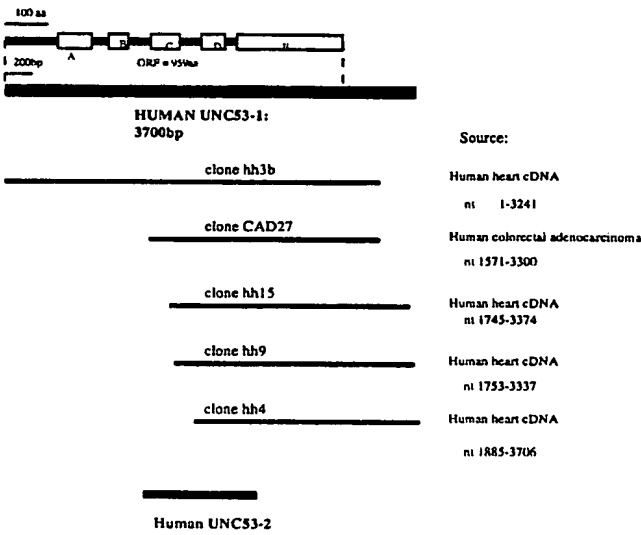


Figure 8

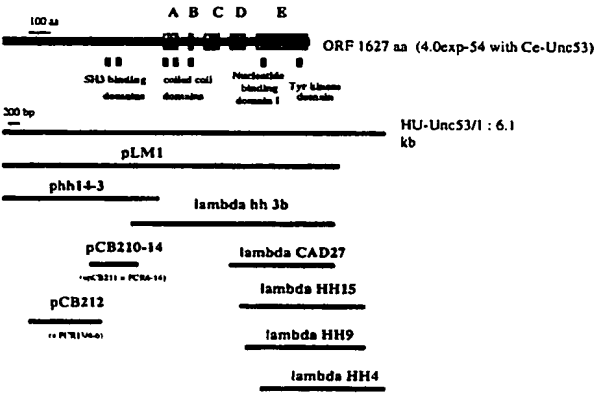


Figure 8a

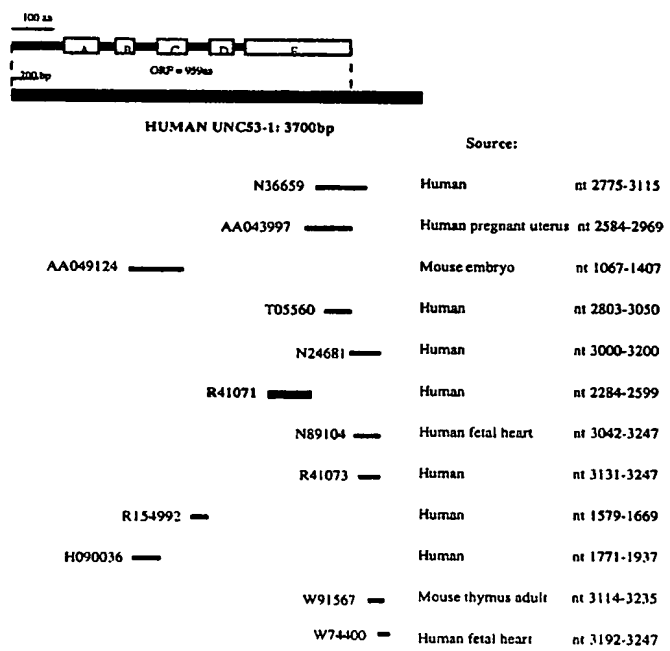


Figure 8b

FIG. 9a

GAAATTCGGGGGAAGGCTGCATTTGGAGTGGAGCTGGGCACTGGGCACTGACAGTAATACAGGACCTCTTCCAGGAAGGGCTCAGG 100
 N S G E E L T V S G S P R A G Q L O S N O R O A N T L P K K G L R
 TACCAGCTTCAGTCCAGGAGGAGACCCAGGAGAGGGGACATTCGCATACCTTGGTGGGCTGCCGAAATCCGATGACGACAGCAGGCTTCCTC 200
 Y O L O S O E E T K E A R H S H T I G G L P E S O O O S E L P S P
 CTGCACCTCCCATGTCTCGAGTGCARAGGGGCACTTACCAACATAGTGGATCCCACTGGCGCCACACGCCAGGAATCACCCGCTCCACAGCATCCC 300
 P A L P N S L S A K G Q L T N I V S P T A T T P R I T A S N S I P
 CACCCACGAGGCGGCCCTTCGAGCTGTACAGCGGCTCCCAATGGGGAGGACCTGTCCCTGGCCGAGAGCCAGGGAATGATTCGGTCAGGATCTTC 400
 T H E A R A F E L Y S G S O N G S T L S L A E R P K G N I R S G S T F
 CGAGACCCACGGAGCATGTTCACGGCTCAGTGTCTCCCTGGCTCCAGTGCCTCCCTACCTACTCTCTGAGCTGAGGAGGAGGATGCAATCTGAGCARR 500
 R O P T O D V H G S V L S L A S S A S S T Y S S A C E E R N O S E O
 homology block A
 TCCGGAGCTTCGTAGGGAATCGAATCATCCAGGAARAGTGGCCACTTGAGCTCTCAGCTTCTGCCAATGCTATCTGGTGGCTGCTTTTGAGCA 600
 I R K L R A E L E S S O E K V A T L T L S A N A N L V A R F E O
 homology block A
 GAGCTCGGTGATATGACATCCCGCTCGGACACTGGCAGAGGCGGCGAGGAGAGGACACTGAGCTGCTGGATTTGCGAGAACCATAGACTTTCTG 700
 S L V N A T S A L A H L A E T A E E K O T E L L O L A E T I O F L
 homology block A
 AAGAAAGAACTCTGAGGCCAGGCGATCATTCAGGGAGCCCTTATGCTCAGAACCAACCAAGAACTCTGGAATCAGAGCAACCACTCTCAG 800
 K K K N S E A O A V I G G A L N A S E T T P K E L A I K R O N S S
 homology block A
 ATAGCATCTCAGGCTCAGCAGCATCACTAGCACTTCAGCATCGGAGCAGCAGGATGCTGATGCGAAAGAGAGAAAGAGATTTGGCTCATGTA 900
 O S I S S L N S I T S H S S I G S S K O A O A K K K K K S V V Y E
 homology block B
 GCTTCGAGATTCCTTACCAAGGCTTCAGTATARAAGGGGCGCCAGTACGCTTCTCTATCTCGGATATAGGAGGATGCTACACCCGACTCTCA 1000
 L A S S F N K A F S I K K G P K S A S S Y S O I E E I A T P O S S
 homology block B
 GCGCCCTCATCTCCCAACATACAGATGGTCTACAGAGACGCTTCACCTCCATCAGTCTCCACCTTGTCTCCGTGGGCACTGATGTACAGCGAGG 1100
 A P S S P K L O H G S T E T A S P S I K S S T L S S V G T O V T E
 GCCCCTGCTACCCAGCCCCCAGCAGTAGGCTGTTCATGCAATGAGGAGGAGGAGCAGAGAGAGGAGGAGGATCTGGAGCTGCGCTCTGAGCATGGGA 1200
 G P A M P A P H T R L F H A N E E E E P E K K E V S E L R S E L V E
 homology block C
 GAGGAATGAGCTTACAGACATCCGCTTGGAGGCCCTCAACTCTGCCCACTTGGATCAGCTTCGGGAGACCATGACACATGCAAGTTGAGGAGT 1300
 K E N K L T O I A L E A L N S A M O L O O L R E T M H N A O L E V
 homology block C
 GACCCTGCTGAAGCAGAGATGACGACCTGAGGCTAGCCCCAGGCCCTCTCAGGCTCCACTCCAGGGCAGGCTCCCTGGATCATCTGCAATATCTTCCC 1400
 O L L K A E N O R L K V A P G P S S G S T P G O V P G S S A L S S
 homology block C
 weak homology in hum1 vs hum2
 CAGCGGCTTCCCTAGGCTGGCACTCAGCATCTCTTGGGCCCCAGCTTGCAGACACAGACTGTACACCATGGAATGGCACTAGTATTTGGTCCAAA 1500
 P R A S L G L A L T H S F G P S L A O T D L S P N O G I S T C G P K
 weak homology in hum1 vs hum2
 GAGGAGGATGACCTTCGGGCTGGTGGTGGGAATGCCCTCCAGCACATCATCAAGGGGCACTGAGCAGCAGGAAATCTTCTTCCCTGGCTGATCAGGCTC 1600
 E E V T L R V V V A N P P O H I I K G O L K O O E F F L G C S K V
 homology block D
 weak homology in hum1 vs hum2
 AGTGGAAAGTTGATGGAAGATGCTGGATGAGCTGTCTTCAGCTGTTCAGGACATATATCTTAAGTGGAGCCAGGCTCTTCCCTGGGATGAGCA 1700
 S G K V O V K A L O E A V F O V F K O V I S K M D P A S T L G L S
 homology block D
 CTGAGTCCATCTCAGCTACAGCATGAGCTAGCTGAAGAGGATGTTGGATGAGAGCCCCCGAGATGCTCTCTTGGCTGGAGGCTCAATACATATC 1800
 T E S I H G Y S I S M V K R V L O A E P P E M P P C A R G V N N I S
 homology block D

FIG. 9a CONTINUED.

AGTC TCCTCCAAAGGTCTGAAGGAGAATCGCGTGCACGCC TGGTGTTTGCAGACCGTAGTCCCAGCACCAGCATGACGACTACATRAAGCTCC TGCTG 1900
V S L K G L E C C V D S L V F E T L I P K P M N O H V I S L L

homology block E - pred nucleotide BD

RAGCACGGGCGCC TCGTCTCTCTC33CCCCAGCGGGCAGGGCAGCACT TACCTGACCACTCGCTTGGCCGAGTACC TGGTGGAGCGCTCTGGCCGTGAGG 2000
K M R A L V L S S P S G T G X Y L T M R L A E V L V E R S G R E

homology block E - pred nucleotide BD

TACAGAGGGGATCG"CAGCAC""ACATGACCCAGCAGTCTTGCAAGGATCTGCAAT TGATATCTTCCARCCCTAGCCARCCAGATAGACGGGAAC 2100
V T E G I V S T F N H M O O S C K O L Q L V L S N L A N O I D R E T

homology block E - pred nucleotide BD

AGCAATTGGGGATGTGCCCC TGGTGATCTATTCGATGACE TAGTGAAGCAGS TCCATCAGTGAG TTGGTCRATGGGCCCC TCACC TGCAAGTATCAT 2200
G I G O V P L V I L L D O L S E R G S I S E L V N G A L T C K Y M

homology block E - pred nucleotide BD

AATGTGCCC ATATA TTAGG TAC CAC CAT CAGC CTG TAARAATGACACCCARCCATGGC TTGCATCTGAGC TTCAGGATGTTCGACTTC ECARACAG 2300
K C P Y I I G T T M O P V K N T P N H G L M L S F A N L T F S N N

homology block E - pred nucleotide BD

TGGAGCAGCCCAATGGCTTCC TGGTCTCTTACCTGAGGAGGAGC TGGTAGAGTACGACAGCGCATCAATGCCAACAGGAGAGCTGCTTCGGGTGCT 2400
V E P A N G F L V A V L R A K L V E S O S O I N A N K E E L L A V L

homology block E - pred nucleotide BD

CGAC TGGGTACCCAGCTGTGGTATCTCTECACACC TTCTTGAGAAGCAGCAGCCTCAGACTTCTCTATCGGCCCTTGTCTCTTCTGTGTGTGCCC 2500
O V V P K L V Y M L M T F L E K H S T S O F L I G P C F F L S E P

homology block E - pred nucleotide BD

ATTGGCAITGAGGACTTCGGGACCTGGTTCATGACC TGTGGARACACTCTATCATTTCCCTATCTACAGGAGGAGGCCAGGA TGGGA TAAGGTCCATG 2600
I G I E O F A T V F I O L V N M S I I P Y L O E G A K O G I K V M

homology block E - pred nucleotide BD

GACAGAARGC TGE TGGGAGGACC CG TGGAA TGGGTCCGGGACACATTCCTTGGCATCAGCCCAACAGACCAATCAAGCTGTACACC TGCECCC 2700
G O K A A V E O P V E V V R D T L P V P S A O O O O S K L Y H L P P

homology block E - pred nucleotide BD

ACCCACCGTGGGCCCTCAGACA TTTCTCTACCTCCCGAGGATAGGACAGTCAAAGCAGCAGCCCAAGTTC TCTGGAC TCAGATCTCTGTATGGCCATG 2800
P T V G P H S : A S P P E O R T V K O S T P S S L O S O P L N A A

homology block E - pred nucleotide BD

CTGC TGAAACTCTAAGAGC TGCCARC TACAT TGAGTCTCCAAGTCAGAGAACA TCC TGGACCCCAACCTTCAGGCACACTTTAAGGGTTCGGCAATC 2900
L L K L O E A A N V I E S P O R E T I L O P N L Q A T L

homology block E - pred nucleotide BD

AC TGTCCCCCCGAC AGCAGACCTCTGCATCAGCTATCTTAGCTCC TCC TC TCCC TC TCC TC TTTCAGAGCAC TGGCT CTCAGCCCCAGGAGGAGA 3000

3' untranslated trailer

ACAGGAGGAGGAGGAGATGAAGGAGG-GGGACAGGTTC TTGGTGC GTACCTTTGAGAACTTCC TAGGAAGGAAATGGTGGG TGGCGT TGGGAC TTG 3100

3' untranslated trailer

TGCCCCCAAAACATTTAC TGCCC "CTCTTAATGAC TT TGGGAAAAGATGATTC TGGGTCTTCCCTTGACTCTTGT TTTCAAT TACAACTTCCTGGG 3200

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3' untranslated trailer

TTAACC 3705

Fig. 9b

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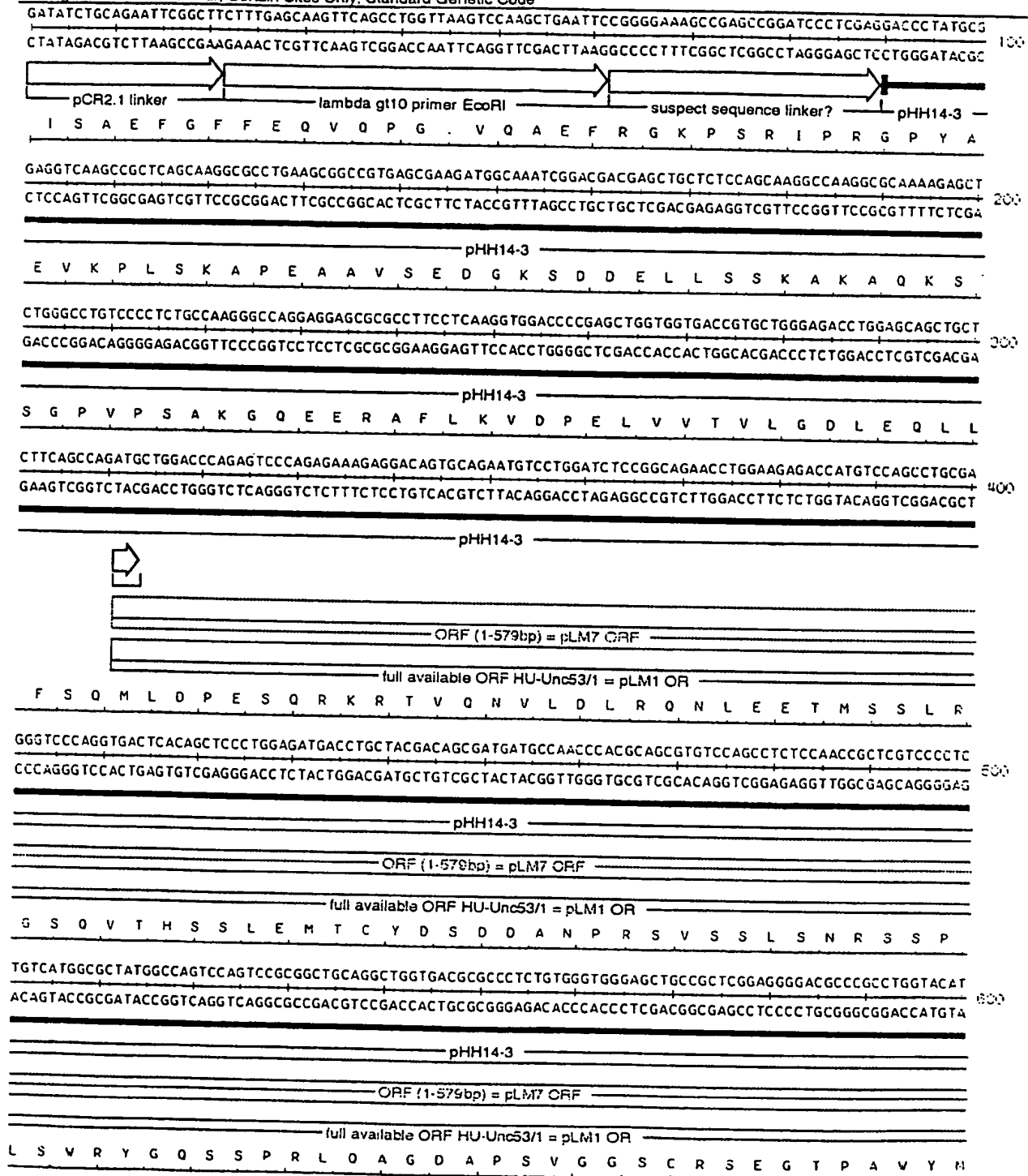
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Settings: Linear, Certain Sites Only, Standard Genetic Code

Page 1

Fig 9 14 pag



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fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 7

GCACGGCGAACGGGCCAC TACTCCCACACCATGCCATGCGCAGCCCCAGCAAGCTCAGCCATATCTCCCGCCTGGAGCTGGTCTGAATCCCTGSACTCG 700
CGTGCCGCTTGCCCGGGTGATGAGGGTGTGGTACGGGTACGGCTCGGGGTCGTTTCGAGTCGGTATAGAGGGCGGACCTCGACCAGCTTAGGGACCTGAGC

————— pHH14-3 —————
————— ORF (1-579bp) = pLM7 ORF —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
H G E R A H Y S H T M P M R S P S K L S H I S R L E L V E S L D S

GATGAGGTGGACCTCAAGTCCGGCTACATGAGCGACAGTGACCTCATGGGCAAGACCATGACGGAGGATGATGACATCACTACCGGCTGGGATGAAAGCA 800
CTACTCCACCTGGAGTTTCAAGCCGATGTACTCGCTGTCACTGGAGTACCCGTTCTGGTACTGCCCTCTACTACTGTAGTGATGGCCGACCTTACTTTCGT

————— pHH14-3 —————
————— ORF (1-579bp) = pLM7 ORF —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
D E V D L K S G Y M S D S D L M G K T M T E D D D I T T G V D E S

GCTCCATCAGTAGTGGACTCAGCGATGCCTCAGACAATCTCAGTTCAGAAGAATTCAATGCCAGCTCCTCACTCAACTCCCTCCCAAGTACTCCCACTGC 900
CGAGGTAGTCATCACCTGAGTCGCTACGGAGTCTGTTAGAGTCAAGTCTTCTTAAGTTACGGTCGAGGAGTGAGTTGAGGGAGGGTTCATGAGGGTGAGC

————— pHH14-3 —————
————— pCB212 —————
————— ORF (1-579bp) = pLM7 ORF —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
S S I S S G L S D A S D N L S S E E F N A S S S L N S L P S T P T A

TTCTCGCAGGAACCTCAACAATAGTGCTACGCACAGACTCAGAGAAGCGCTCACTGGCAGAAAGTGGGCTGAGCTGGTTAGTGAATCAGAGGAGAAAGCC 1000
AAGAGCGTCTTGAGTTGTTATCACGATGCGGTGCTGAGTCTCTTCGCGAGTGACCGTCTTTCACCCGACTCGACCAATCACTTAGTCTCTCTTTGGG

————— pHH14-3 —————
————— pCB212 —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
S R R N S T I V L R T D S E K R S L A E S G L S W F S E S E E I A

CCTAAAAAAGTGGAGTACGACAGTGGTAGCCTGAAGATGGAACCTGGGACTTCTAAGTGGCGGAGGGAGCGGCTGAGAGCTGTGATGATTCATCCAAAGG 1100
GGATTTTTTGACCTCATGCTGTCAACATCGGACTTCTACCTTGACCCCTGAAGATTCACCGCTCCCTCGCCGGACTCTCGACACTACTAAGTAGGTTCC

————— pHH14-3 —————
————— pCB212 —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
P K K L E Y D S G S L K M E P G T S K V R R E R P E S C D D S S I

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fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 1

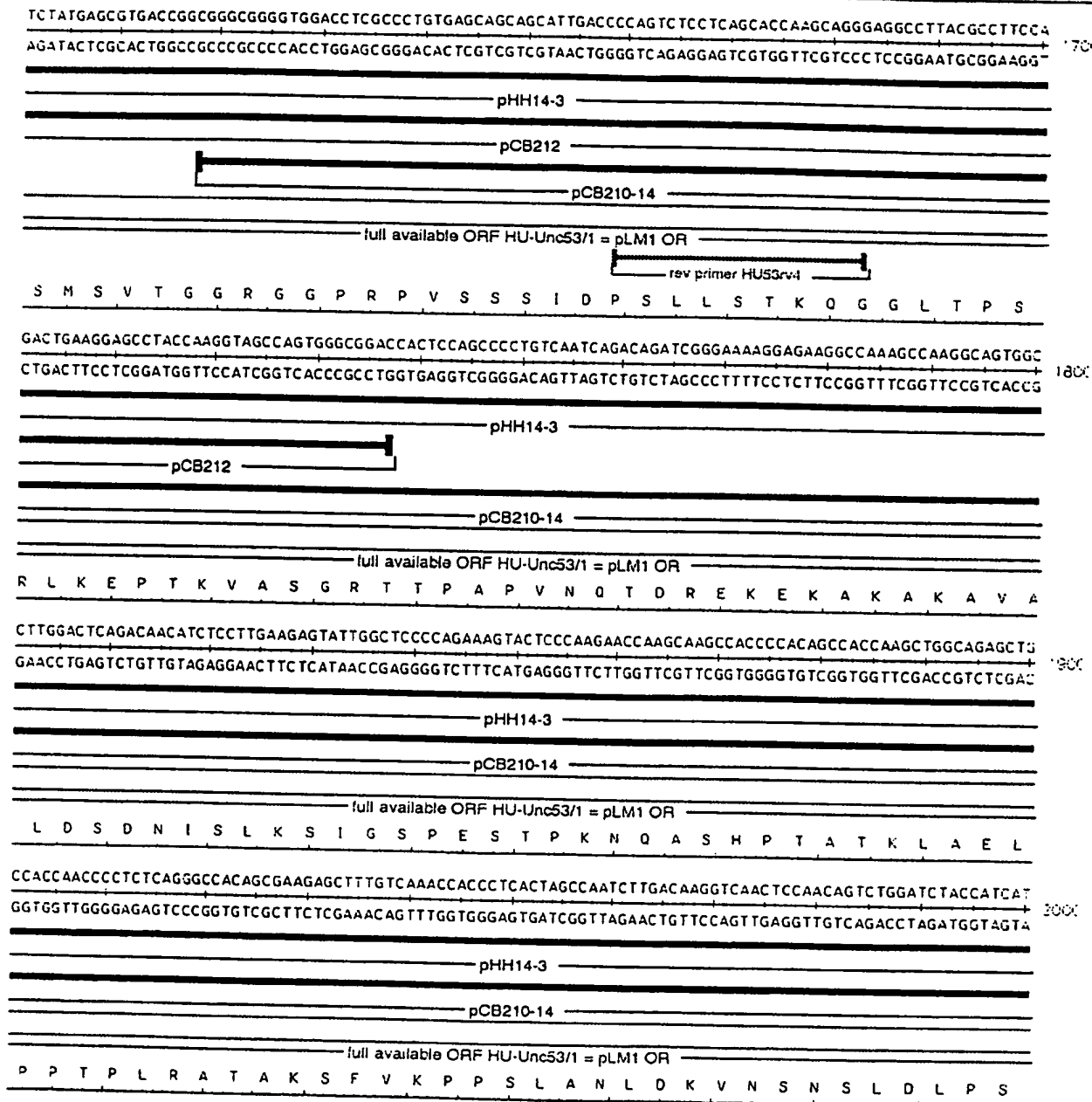
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CACCTCTTGACTTTTTTCGGGTAGTCGGACCCGGTGGGACCAAGGGACTTCTTCCCGTTCTGGGGTGGACACCGACATTGAAGGGGGTAGTGAGTGTGTCT 120

————— pHH14-3 —————
————— pCB212 —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
G G E L K K P I S L G H P G S L K K G K T P P V A V T S P I T H T A
CCAGAGTGGCCTCAAAGTCGCAGGCAAACTGAGGGCAAGCTACAGACAAGGGTAAGCTTGCACTGAAGAATACTGGGCTCCAACGCTCCCTCTCTGAT 130
GGTCTCACGGGAGTTTCAGCGTCCGTTTGGACTCCCGTTTCGATGCTGTTCCTTCGAACGTCACCTTCTATGACCCGAGGTTCGAGGAGGAGACTA
————— pHH14-3 —————
————— pCB212 —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
O S A L K V A G K P E G K A T D K G K L A V K N T G L O R S S S D
GCTGGTCGGGACCGCCTGAGTGATGCTAAGAAGCCCCCTCGGGCATTGCTCGCCCCCTCCACTTCGGGATCCTTTGGCTACAAGAAGCCTCCCTCTGCCA 140
CGACCAGCCCTGGCGGACTCACTACGATTCTTCGGGGGAGCCCGTAACGAGCGGGGAGGTGAAGCCCTAGGAAACCGATGTTCTTCGGAGGAGGACGGT
————— pHH14-3 —————
————— pCB212 —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
A G R D R L S D A K K P P S G I A R P S T S G S F G Y K K P P P A
CAGGCACAGCCACTGTATGCAAACTGGTGGTTCAGCCACTCTCAGCAAGATCCAGAAGTCTCAGGCATCCCTGTCAAGCCAGTAAATGGGCGCAAGAC 150
GTCCGTGTGGGTGACAGTACGTTTGACCACCAAGTCGGTGAGAGTCTGTAGGTCTTCAGGAGTCCGTAGGGACAGTTCGGTCATTACCCGCGTCTCT
————— pHH14-3 —————
————— pCB212 —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
T G T A T V M O T G G S A T L S K I O K S S G I P V K P V N G R F T
TASCTTAGATGTTTCCAACAGTGCAGAGCCAGGATTCTTGGCTCCTGGAGCCCGTTCTAACATCCAGTACCGCAGCCTGCCCCGGCCAGCCAAGTCAAGT 160
ATCGAATCTACAAAGGTTGTCAGTCTCGGTCTTAAGGACCGAGGACCTCGGGCAAGATTGTAGGTTCATGGCGTCGGACGGGGCCGGTTCAGTTCA
————— pHH14-3 —————
————— pCB212 —————
————— full available ORF HU-Unc53/1 = pLM1 OR —————
S L D V S N S A E P G F L A P G A R S N I Q Y R S L P R P A K S S

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fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 4



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fig. 19. Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 5

CCAGTGATACCACCCATGCTTCAAAGGTCCCAGATCTGCATGCTACAAGCTCAGCATCTGGGGGCCCTCTCCCTTCTGCTTACCCCCAGTCCGGCACC
GGTCACTATGGTGGGTACGAAGTTTCCAGGGTCTAGACGTACGATGTTGAGTCGTAGACCCCGGAGAGGGAAGGACGAAGTGGGGGTCAGGCCGTGG 210

pHH14-3

pCB210-14

full available ORF HU-Unc53/1 = pLM1 OR

S S D T T H A S K V P D L H A T S S A S G G P L P S C F T P S P A P

CATCTCAATATTAACTCAGCCAGCTTCTCCAGGGCCTGGAGCTAATGAGTGGTTTCAGTGTGCCAAAAGAGACCCGCATGTACCCCAAACCTCAGGC
GTAGGAGTTATAATTGAGTCGGTCAAGAGGGTCCCGGACCTCGATTACTACCAAAGTCACACGGTTTTCCTGGGCGTACATGGGGTTTGAGAGTCCG 220

pHH14-3

pCB210-14

full available ORF HU-Unc53/1 = pLM1 OR

I L N I N S A S F S Q G L E L M S G F S V P K E T R M Y P K L S G

CTGCACAGGAGCATGGAGTCCCTCCAGATGCCAATGAGCCTCCCAAGTGCCTTCCCAGCAGTACTCCCGTCCCCACCCACCTGCTCCCCCTGTGTCTC
GACGTGTCTCGTACC TCAGGGAGGTCTACGGTTACTCGGAGGGGTACGGAAGGGTCTGTCATGAGGGCAGGGGTGGGGTGGACGAGGGGGACGACGAG 230

pHH14-3

pCB210-14

full available ORF HU-Unc53/1 = pLM1 OR

L H R S M E S L Q M P M S L P S A F P S S T P V P T P P A P A A

CCACAGAAGAAGAGACGGAAGAGCTGACTTGGAGTGGGAAGCCCCAGAGCTGGGCAACTGGACAGTAATCAGCGGGATCGGAACACTCTTCCCAAGAAAGG
GGTGTCTTCTCTCTGCCTTCTCGACTGAACCTACCTTCGGGGTCTCGACCCGTGACCTGTCAATTAGTCGCCCTAGCCTTGTGAGAAGGGTTCTTTCC 240

pHH14-3

pHH3b

pCB210-14

full available ORF HU-Unc53/1 = pLM1 OR

rev primer HU53rv3

rev primer HU53rv2

peptide B72828H

P T E E E T E E L T V S G S P R A G Q L D S N Q R D R N T L P K K G

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fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 4

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CGAGTCCATGGTTCGAAGTCAGGGTCTCTCTGGTTCCTCTCCGCTGTAAGGGTATGGTAACCCCGACGGACTTAGGCTACTGGTCAGTCTCGACGGG

pHH14-3

pHH3b

rev primer HU53rv1

full available ORF HU-Unc53/1 = pLM1 OR

L R Y Q L Q S O E E T K E R R H S H T I G G L P E S D D Q S E L P

TCTCCCCCTGCACCTCCCATGTCTCTGAGTGCAGAGGCGCAACTTACCAACATAGTGAGTCCCAGTCCGCGCCACCACGCCAAGAAATCACCCGCTCCAACA 260
AGAGGGGGACGTGAAGGGTACAGAGACTCACGTTTCCCGGTGAATGGTTGTATCACTCAGGGTGACGCCGGTGGTGCCTTCTAGTGGGCGAGGTTGT

pHH14-3

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S P P A L P M S L S A K G Q L T N I V S P T A A T T P R I T R S N

GCATCCCCACCCACGAGGCGGCC TTCGAGCTGTACAGCGGCTCCCAATGGGGAGCACCTGTCCCTGGCCGAGAGACCCAAGGAATGATTCGGTCAGG 270
CGTAGGGGTGGTGTCTCCGCCGAAGCTCGACATGTGCCGAGGGTTTACCCTCGTGGGACAGGGACCGGCTCTCTGGGTTCCCTTACTAAGCCAGTCC

pHH14-3

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S I P T H E A A F E L Y S G S Q M G S T L S L A E R P K G M I R S G

ATCCTTCCGAGACCCACGGACGATGTTACGGCTCAGTGCTGTCCCTGGCCTCCAGTGCTCTCCACCTACTCCTCAGCTGAGGAGAGGATGCAATCT 280
TAGGAAGGCTCTGGGGTGCTGTCTACAAGTGCCGAGTCACGACAGGGACCGGAGGTACGGAGGAGGTGGATGAGGAGTCGACTCCTCTCTACGTTAGA

pHH14-3

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

S F R D P T D D V H G S V L S L A S S A S S T Y S S A E E R M Q S

GAGCAATCCGGAAGCTTCGTAGGGAAC TGAATCATCCCAGGAAAAAGTGGCCACCTTGACGTCTCAGCTTTCTGCCAATGCTAATCTGGTGGCTGCTT 290
CTCGTTAGGCCCTTCGAAGCATCCCTTGACCTTAGTAGGGTCTTTTACCAGGTGGAACGCAGAGTCGAAAGACGGTTACGATTAGACCACCGACGAA

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

E O I R K L R R E L E S S O E K V A T L T S O L S A N A N L V A A

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fig Hu-Unc53/1 seq (1>6013) Site and Sequence

Fig 95

Page 7

TTGAGCAGAGCCTGGTGAATATGACATCCCGCCTGCGACACCTGGCAGAGACGGCCGAGGAGAAGGACACTGAGCTGCTGGATTTCGAGAAACCATAGA 300
AACTCGTCTCGGACCACCTTACTGTAGGGCGGACGCTGTGGACCGTCTCTGCCGGCTCCCTTCCCTGTGACTCGACGACCTAAACGCTCTTTGGTATCT
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
F E Q S L V N N T S R L R H L A E T A E E K D T E L L D L R E T I D
CTTTCGAAGAAAAGAACCTGAGGCCAGGCGATCATTCAGGAGGCCCTTAATGCC TCAGAAACCACACCCAAAGAAC TTCGGATCAAGAGACAAAAC 310
GAAAGACTTC TTTTCTTGAGACTCCGGGTCCGT CAGTAAGTCCCTCGGGAATTACGGAGTCTTTGGTGTGGGTTTCTTGAAGCC TAGTCTCTGTTTGT
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
F L K K K N S E A Q A V I Q G A L N A S E T T P K E L R I K R Q N
TCCTCAGATAGCATCTCAAGCCTCAACAGCATCAC TAGCCATTCAGCATCGGCAGCAGCAAGGATGCTGATGCGAAAAAGAAGAAAAAAGAGTTGGG 320
AGGAGTCTATCGTAGAGTTCGGAGTTGTCGTAGTGATCGGTAAGGTCGTAGCCGTCGTCGTTCC TACGACTACGCTTTTTCTCTTTTTTTCTCAACCC
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
S S D S I S S L N S I T S H S S I G S S K D A D A K K K K K K S V
TCTATGAGCTTCGAAGTTCTTCAACAAAGCGTTCAGTATAAAAAAGGGGCCAAGTCAGCTTCTCATACTCGGATATAGAGGAGATTGCTACACCCGA 330
AGATACTCGAAGCTTCAAGGAAGTTGTTTCGCAAGTCATATTTTTCCCGGGTTCAGTCGAAGGAGTATGAGCCTATATCTCCTCAACGATGTGGGCT
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
V Y E L R S S F N K A F S I K K G P K S A S S Y S D I E E I A T P D
CTCTTCAGCCCCCTCATCCCCAAACTACAGCATGGTTC TACAGAGACTGCTTCACCCCTCCATCAAGTCTCCACCTTGCTCCTCCGTGGGACTGATGTC 340
GAGAAGTCGGGGGAGTAGGGGGTTTGATGTCTACCAAGATGTCTCTGACGAAGTGGGAGGTAGTTCAGGAGGTGGAACAGGAGGCACCCGTGACTACAG
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
S S A P S S P K L Q H G S T E T A S P S I K S S T L S S V S T D V

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file: hu-Unc53/1 seq (1 > 6013) Site and Sequence

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Fig 9b

ACCG .GCCCTGCTCACCAGCCCCACACTAGGCTGTTCATGCAATGAGGAGGAGGAGCCAGAGAAGAAGGAGGTATCGGAGCTGCGCTCTGAGG 3500
TGGCTCCCGGGACGAGTGGGTCGGGGGGTGTGATCCGACAAGGTACGTTTACTCCTCCTCCTCGGTCTCTTCTTCTCCATAGCCTCGACGCGAGACTCG
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
T E G P A H P A P H T R L F H A N E E E E P E K K E V S E L R S E
TATGGGAGAAGGAAATGAAGCTTACAGACATCCGCTTGGAGGCCCTCAACTCTGCCACCAACTGGATCAGCTTCGGGAGACCATGCACAACATGCAGTT 3600
ATACCCCTCTTCTTTACTTCGAATGTCTGTAGGCGAACCCTCCGGGAGTGTAGAGCGGGTGGTTGACCTAGTCGAAGCCCTCTGGTACGTGTTGTACGTCAA
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- peptide B72527H -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
----- U3 ORF = pLM5 ORF -----
L V E K E M K L T D I R L E A L N S A H Q L D Q L R E T M H N M Q L
GGAGGTGGACCTGCTGAAAGCAGAGAATGACCGAC TGAAGGTAGCCCCAGGCCCTCATCAGGC TCCACTCCAGGGCAGGTCCCTGGATCATCTGCATTA 3700
CCTCCACCTGGACGACTTTCGTCCTTACTGGCTGACTTCCATCGGGTCCGGGGAGTAGTCCGAGGTGAGGTCCCGTCCAGGGACCTAGTAGACGTAAT
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
----- U3 ORF = pLM5 ORF -----
E V D L L K A E N D R L K V A P G P S S G S T P G Q V P G S S A L
TCTTCCCCAGCCGCTCCCTAGGCCCTGGCACTCACCCATTCTTCCGCCCCAGTCTTGACAGACAGACCTGTACCCATGGATGGCATCAGTACTTGTG 3800
AGAAGGGGTGCGGCGAGGGATCCGGACCGTGAGTGGTAAGGAAGCCGGGGTCAGAAGCTCTGTGCTGGACAGTGGGTACC TACCGTAGTCATGAACAC
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
----- U3 ORF = pLM5 ORF -----
S S P R R S L G L A L T H S F G P S L A D T O L S P M D G I S T C

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file 'u-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 9

GTCCAAAGGAGGAAGTGACCTCCGGGTGGTGGTGAGGATGCCCCCGCAGCACATCATCAAAGGGGACTTGAAGCAGCAGGAATTCTTCCTGGGCTGTAG
CAGGTTTCTCTCTTCACTGGGAGGCCACCACCACCTCTACGGGGGCGTGTGTAGTAGTTTCCCTGAAC TTCGTGTCCTTAAGAAGGACCCGACATC

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

G P K E E V T L R V V V R M P P O H I I K G D L K Q Q E F F L G C S

CAAGGTCAGTGGAAAAGTTGACTGGAAGATGCTGGATGAAGCTGTTTTCCAAGTGTTCAGGACTATATTTCTAAAATGGACCCAGCCTCTACCCTGGGA
GTTCCAGTCACCTTTTCAACTGACCTTCTACGACCTACTTCGACAAAAGGTTACAAGTTCC TGATATAAAGATTTTACC TGGGTGGGAGATGGGACCCCT

U2 ORF = pCB251 ORF

pHH3b

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

K V S G K V D V K M L D E A V F Q V F K D Y I S K M D P A S T L G

CTAAGCACTGAGTCCATCCATGGCTACAGCATCAGCCACGTGAAACGAGTGTGGATGCAGAGCCCCCGAGATGCCTCCTTGCCGTCGAGGTGTCAATA
GATTCGTGACTCAGGTAGGTACCGATGTCGTAGTCGGTGACATTTGCTCACAACCTACGTCTCGGGGGGCTCTACGGAGGAACGGCAGCTCCACAGTTAT

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

L S T E S I H G Y S I S H V K R V L D A E P P E M P P C R R G V N

ACATATCAGTCTCCCTCAAAGGTCTGAAGGAGAAAATGCGTCGACAGCCTGGTGTTCGAGACGCTGATCCCCAAGCCGATGATGCAGCACTACATAAGCCT
TGATATAGTCAGAGGGAGTTTCAGACTTCTCTTTACGCAGCTGTCGGACCAACAAGCTCTGCGACTAGGGGTTCGGCTACTACGTCGTGTGTATTGGGA

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

H I S V S L V G L K E K C V D S L V F E T L I P K P M M Q H Y I S L

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fig. 14-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 10

CCCTCTGAGCACCAGGCGCCTCGTCCTCTCGGGCCCCAGCGGCACGGGCAAGACCTACCTGACCAATCGCTTGGCCGAGTACCTGCTGGAGCGCTCTGGC
GGACGACTTCGTGGCCGCGGAGCAGGAGAGCCCCGGGTCGCCGTGCCGTCTTGGATGGACTGGTTAGCGAACCGGCTCATGGACCACTCGCGAGACCG
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- U4 ORF = pCB201 ORF -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
----- U3 ORF = pLM5 ORF -----
----- pHH15 -----
L L K H R R L V L S G P S G T G K T Y L T N R L A E Y L V E R S G
CGTGAGGTACAGAGGGCATCGTCAGCACCTTCAACATGCACCAGCAGTCTTGCAAGGATCTGCAACTGTATCTTTCCAACCTAGCCAACCCAGATAGACC
GCACTCCAGTGTCTCCCGTAGCAGTCGTGGAAGTTGTACGTGGTCGTCAGAACGTTCTTAGACGTTGACATAGAAAGGTTGGATCGGTTGGTCTATCTGG
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- U4 ORF = pCB201 ORF -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
----- U3 ORF = pLM5 ORF -----
----- pHH15 -----
R E V T E G I V S T F N M H Q Q S C K D L Q L Y L S N L A N Q I D
GGGAAACAGGAATTGGGGATGTGCCCTGGTGATTCTATTGGATGACCTGAGTGAAGCAGGC TCCATCAGTGAGTTGGTCAATGGGGCCCTCACTGCA
CCCTTTGTCTTAACCCCTACACGGGGACCACTAAGATAACCTACTGGACTCACTTCGTCCGAGGTAGTCACTCAACCAGTTACCCCGGGAGTGGACGT
----- U2 ORF = pCB251 ORF -----
----- pHH3b -----
----- U4 ORF = pCB201 ORF -----
----- full available ORF HU-Unc53/1 = pLM1 OR -----
----- U3 ORF = pLM5 ORF -----
----- pHH15 -----
R E T G I G D V P L V I L L D D L S E A G S I S E L V N G A L T C I

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fig. ⁴Hu-Unc53/1 seq (1 > 8013) Site and Sequence

Fig 9b

Page 17

GTATCA:AAATGTCCCTATATTATAGGTACCACCAATCAGCC TGTAAAAATGACACCAACCATGGCTTGCAC TTGAGCTTCAGGATGTTGACCTTCCTC
CATAGTATTTACAGGGATATAATATCCATGGTGGTTAGTCGGACATTTTACTGTGGGTGGTACCGAAGCTGAAC TCGAAGTCCTACAAC TGGAAAGAGG 460

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

peptide B72626H

Y H K C P Y I I G T T N Q P V K M T P N H G L H L S F R M L T F S

AACAACGTGGAGCCAGCCAATGGCTTCCTGGTTCGTTACCTGAGGAGGAAGCTGGTAGAGTCAGACAGCGACATCAATGCCAACAAGGAAGAGC TGCTTC
TTGTTGCACCTCGGTCGGTTACCGAAGGACCAAGCAATGGACTCCTCCTTCGACCATCTCAGTCTGTCGCTGTAGTTACGGTTGTTCTCTTCGACGAAG 470

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

N N V E P A N G F L V R Y L R R K L V E S D S D I N A N K E E L L

GGGTGCTCGACTGGGTACCCAAGCTGTGGTATCATCTCCACACCTTCCTTGAGAAGCACAGCACCTCAGACTTCCTCATCGGCCCTTGCTTCTTTCTGTG
CCCACGAGCTGACCCATGGGTTGACACCATAGTAGAGGTGTGGAAGGAACCTTCGTCGTCGTGGAGTCTGAAGGAGTAGCCGGGAACGAAGAAAGACAG 480

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

R V L D V V P K L V Y H L H T F L E K H S T S D F L : G P C F F L S

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fig. HU-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 12

GTSTCCCATTTGGCATTGAGGACTTCCGGACCTGGTTCATTGACCTGTGGAACAACCTCTATCATTCCTTATCTACAGGAAGGAGCCAAGGATGGGATAAAG
CAGAGGGTAACCGTAACCTCTGAAGGCCCTGGACCAAGTAAGTGGACACCTTGTGAGATAGTAAGGGATAGATGTCTTCTCGGGTCTTACCTATTTCT

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

C P I G I E D F R T V F I D L V N N S I I P Y L Q E G A K D G I I

GTCCATGGACAGAAAGCTGCTTGGGAGGACCCAGTGGAAATGGGTCCGGGACACACTTCCCTGGCCATCAGCCCAACAAGACCAATCAAAGCTGTACCACT
CAGGTACCTGTCTTTCGACGAACCTCTCGGGTCACCTTACCCAGGCCCTGTGTGAAGGGACCGGTAGTCGGGTGTCTGGTTAGTTTCGACATGGTGG

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

V H S O K A A V E D P V E W V R D T L P V P S A O Q D O S K L Y H

TGCCCCCACCACCGTGGGCCCTCACAGCATTGCCTCACCTCCCGAGGATAGGACAGTCAAAGACAGCACCCCAAGTTCTCTGGACTCAGATCCTCTGA
ACGGGGGTGGGTGGCACCCGGGAGTGTGTAACGGAGTGGAGGGCTCTATCTGTGAGTTTCTGTCGTGGGGTTCAAGAGACCTGAGTCTAGSAGACTA

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

L P P P T V G P H S I A S P P E D R T V K D S T P S S L D S O P L I

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fig. 4u-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 13

GGCCATGCTGCTGAAACTTCAAGAAGCTGCCAATACATGAGTCTCCAGATCGAGAAACCATCTGGACCCCAACCTTCAGGCAACACTTTAAGGGTTC
CCGGTACGACGACTTTGAAGTCTTCGACGGTGTGTAATCAGAGGTCTAGCTCTTGGTAGGACCTGGGGTTGGAAGTCCGTGTGAAATTCCTCAAG 520

U2 ORF = pCB251 ORF

pHH3b

U4 ORF = pCB201 ORF

full available ORF HU-Unc53/1 = pLM1 OR

U3 ORF = pLM5 ORF

pHH15

peptide B72625H

A M L L K L Q E A A N Y I E S P D R E T I L D P N L Q A T L . G F

GGCAATCACTGTACCCCCGGACAGCAGAAGCTGGCATCAGCTATCTTAGCTCCTCCTCTCCCTCTCCTCTTCAGAGCACTGGCTCTCCAGCCCCAG
CCGTTAGTGACAGTGGGGGCTGTCGCTTTCGACCGTAGTCGATAGAATCAGGAGGAGAGGGGAGAGGAGAAAGTCTCGTGACCGAGAGGTCGGGGTC 530

pHH3b

pHH15

G N H C H P R T A E R V H Q L S . L L L S P L L F Q S T G S P A P

GAGGAGAACAGGAGGAGGAGGAGATGAAAGAGSAGGACAGGTTCTTGGTGCTGTACCTTTGAGAACTTCCTAGGAAGGAATGGTGGGGTGGCGTTTGG
CTCCTCTTGCTCCCTCCTCCTCTACTTTCTCCTCCCTGTCCAAGAACCACGACATGGAACTCTTGAAGGATCCTTCTTACCACCCACCGCAAAC 540

pHH3b

pHH15

G G E Q E G G G D E R G G T G S V C C T F E N F L G R N G G V A F G

GAACTTGTCCTCCCTAAACACATTTACTGGCTCCTCTAATGACTTTGGGGAAGATGATTCTGGGTCTTTCCTTGACTTCTTGTTCATTACAAAC
CTTGAACACGGGGGATTGTGTAAATGACCGGAGGAGATTACTGAAACCCCTTTCTACTAAGACCCAGAAAGGGAACGAAGAACAAGTTAATGTTT 550

pHH3b

pHH15

N L C P L N T F T G L L . L V G K D D S G S F P . L L V S I T H

TCTGGGCTTTCTGGGAGGGGTTCAAGAAACATCAAAACACGACGAGTTCTAAATGATTCTCACAAGCAACCTGAGAGAGACAGTCTTGTGAGGG
AGGACCCGAAAGACCCCTCCCAAGTCTTTGTAGTTTGTGACGTCGTCAGGATTACTAAGAGTGTTCGTTGGGACTCTCTGTCAGAACAC TCCC 560

pHH3b

pHH15

S V A F V G G V G K T S K H C S S S . M I L T S N P E R D S L V P

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fig Hu-Unc53/1 seq (1 > 6013) Site and Sequence

Fig 9b

Page 4

AGATCTGGGGGAGGCAGGAAGCTCCTCAGATTTTCACAGACCCTTCCCAATCCATCACCCTGCCAACAACTCTCCCCCAGAGATCTGGCTGGAGC 570
TCTAGACCCCTCCGTCCTTCGAGGAGTCTAAAAGAGTGCTGGGAAGGGTTAAGGTAGTGGTGACGGTTGTTGAGGAGGGGCTCTAGACCGACCTCC 580
pHH15
E I V G R Q E A P Q I F S Q T L P N S I T T A N N S S P R D L A G A
CCAGAAAAAGAAGCATGTGGTTAAAAAATGTTTAAATCAATCTGTAAAAGGTAAAAATGAAAAACAAAAACAAAGCAACAAACAAAAACAATGGAAA 590
GGTCTTTTCTTCGTACACCAATTTTTACAAATTTAGTTAGACATTTCCATTTTACTTTTGTGTTTGTTCGTTTGTGTTTGTGTTTGTGTTACCTTT 600
O K K K H V V . K M F K S I C K R . K . K N K N K Q T N K K Q W I
AGATGAAGCTGGAGAGAGAGGAACCAAGTTGCCAAGGTAGAGAGCTGCCCGCTCCTGCCCTCTGGATGACATAGGGGACATCAACAAGACGGCTGCCAAC 610
TCTACTTCGACCTCTCTCCTTGGTCAACGGTTCCATCTCTCGACGGGCGAGGACGGGAGACCTACTGTATCCCTGTAGTTGTCTGCCGACGGTTGG 620
R . S W R E R N Q L P R . R A A R S C P L D D I G D I N K T A A N
TGAGAAGTCACCAAAACCACAAAATAACCTTACAGCCTTCAGGGAAGACTACCAGCTCTGTCTTTCTACCCTCTAATTTAACAATGCACCGGAATTCAG 630
ACTCTTCAGTGGTTTGGTGTGTTTATTGGAATGTCGGAAGTCCCTTCTGATGGTCGAGACAGAAAGATGGGAGATTAAATTGTTACGTGGCCTTAAGTC 640
linker?
L R S H Q T T K I T L Q P S G K D Y Q L C L S T L . F N N A P E F S
CTTGGACTTAACC 6013
GAACCTGAATTGG
linker?
L D L T

FIG. 10.

GGCACGAGGCATCCTCTGTGGGCACCGAGGTCACCGAGACCCCTGCTCATTCAGTCCCCCACACTAGACT 70
linker ? open reading frame

R H E A S S V G T E V T E T P A H S V P H T R L
GTTCCAAGCCAATGAAGAGGAGGAGCCAGAGAAGAAGGAGGTATCAGAACTGCGCTCTGAACTATGGGAA 140
open reading frame

F Q A N E E E E P E K K E V S E L R S E L V E
AAAGAGATGAAGCTCACGGATATCCGGTTGGAGGCCCTCAACTCTGCCCAACCAGCTGGACCAGCTTCGGG 210
open reading frame

K E M K L T D I R L E A L N S A H O L D O L R
AGACCATGCACAATATGCAGTTGGAGGTGGACCTGCTGAAAGCAGAGAATGACCGGCTGAAGGTGCCCC 280
open reading frame

E T M H N M Q L E V D L L K A E N D R L K V A P
CGGCCCCCTCTCAGGCTGCACTCCAGGGCAGGTCCTGGGTCATCGGCTCTGTCGTCCTCCCTCGACGTTC 350
open reading frame

G P S S G C T P G O V P G S S A L S S P R S
CTGGGCCTTGCACTCAGCCATCCTTTTCAGTCTTAGTCTCACAGACACAGACCTCTCACCCATGGAAGGCA 420
open reading frame

L G L A L S H P F S P S L T D T G L S P M C G
TCAGCACCTGTGGTTCAAAGGAAGAGGTGACCCTGCGGGTGGTGGTCCGGATGCCGCCCCAGCACATCAT 490
open reading frame

I S T C G S K E E V T L R V V V R M P P D H !
CAAAGGGGACTTAAAGCAGCAGGAGTTCTTCTGGGTTGCAGCAAGGTGAGTGGCAAGTTGACTGGAG 560
open reading frame

K G D L K Q Q E F F L G C S K V S G K V D . K
ATGCTGGATGAAGCCGTTTTCCAAGTGTTCAGGACTACATTTCTAAAAAGGAGCCAGCCCTCAAGCTGG 630
open reading frame

M L D E A V F Q V F . D Y I S K M D F A S T L
GACTGAGCACTGAGTCCATACATGGCTATAGCCTCAGCCACGTGAACGAGTGCTGGATGCTGAGGCCCT 700
open reading frame

G L S T E S I H G Y S L S H V K R V L D A E F P

FIG. 10 CONTINUED.

AGAGATGCCCTCCTTGCCGCCGAGGTGTCAATAACATATCAGTCGCTCTCAAAGGTCTGAAAGAGAAGTGT 770
open reading frame

E M P P C R R G V N N I S V A L K G L K E K C
GTCGACAGCCTGGTGTTCGAGACGCTTATCCCAAGCCCATGATGCAGCACTACATCAGCCTCCTGCTCA 840
open reading frame

V D S L V F E T L I P K P M M O H Y I S L L L
AGCACCGGCGCC TGGTGCTCTCCGGCCCCAGTGGCACCGGCAAGACCTACTTGACCAATCGGCTAGCCGA 910
open reading frame

K H R R L V L S G P S G T G K T Y L T N P L A E
GTACCTGGTGGAGCGCTCCGGCCGCGAGGTACGGATGGCATCGTCAGCACTTTCAACATGCACCAGCAG 980
open reading frame

Y L V E R S G R E V T D G I V S T F N M H Q Q
TCTTGCAAGGATCTGCAACTGTACCTCTCCAACCTAGCCAACCAGATAGACCGGGAAACAGGGATAGGGG 1050
open reading frame

S C K D L Q L Y L S N L A N O I D R E T G I G
ATGTGCCCTTGGTGATCCTCCTGGATGATCTGAGTGAAGCAGGCTCCATCAGTGAGCTGGTCAATGGGGC 1120
open reading frame

D V P L V I L L D D L S E A G S I S E L V N G A
CCTCACCTGCAAGTATCACAAATGTCCCTACATTATAGGTACCACCAATCAGCCTGTAAAAATGACACCC 1190
open reading frame

L T C K Y H K C P Y I I G T T N O P / K M T P
AACCATGGCTTGCACITGAGCTTCAGGATGCTGACCTTCTCGAACAATGTGGAACCAGCCCATGGCTTTTC 1260
open reading frame

N H G L H L S F R M L T F S N N V E P A N G F
TGGTCCGTTACCTGCGGAGGAAGTTGGTAGAGTCAGACAGTGACGTCAATGCTAACAGGGAAGAGCTTCT 1330
open reading frame

L V R Y L R R K L V E S D S D V N A N I E E L L
TCGGGTGCTGGACTGGGTGCCCAAGCTGTGGTATCACCTCCACACCTTCTGGAGAGGCAAGCACCTCG 1400
open reading frame

R V L D V V P L V V H L H T F L E . H S T S

FIG. 10 CONTINUED.

GACTTCCTCATTGGCCCTTGCTTCTTCCTGTCCCTGTCCTTGGCATCGAGGACTTCCGGACCTGGTTCA 1470
open reading frame

D F L I G P C F F L S C P I G I E D F R T V F
TTGACCTGTGGAACAATCCATCATCCCCATCTACAGGAAGGAGCCAAGGATGGGATCAAGGTTTCATGG 1540
open reading frame

I D L V N N S I I P Y L Q E G A K D G I K V H G
ACAGAAAGCTGCTTGGGAAGACCCGGTGAATGGGTCCGAGACACTCTTCCCTGGCCGTCGGCCCAACAA 1610
open reading frame

Q K A A V E D P V E V V R D T L P V P S A Q Q
GACCAATCAAAGCTCTACCACCTGCCCCCGCTTCTGTGGGCCCCACAGCACTGCCTCACCCCCGGAGG 1680
open reading frame

D Q S K L Y H L P P P S V G P H S T A S P P E
ACAGGACAGTCAAAGACAGCACTCCAACTCCCTCGACTCAGATCCCTTGATGGCCATGCTACTGAAACT 1750
open reading frame

D R T V K D S T P N S L D S D P L H A M L L K L
CCAAGAAGCTGCCAACTACATTGAGTCACCAGATCGAGAGACTATCCTGGACCCCAACCTCCAGGCGACA 1820
open reading frame

Q E A A N Y I E S P D R E T I L D P N L Q A T
CTCTGAGGGCCCGGCAGTCACTGTCAACCTGGAGGGCAGAAGGCTGGCTTCAGCATCATTAGCTCTCCTC 1890
3' untranslated

L . G P G S H C H P G G Q K A G F S I I S S P
TGCCCTCTTCCTTCATAGCTCTGGCTCACCAGCCTCGCCAAGAGAACAGGAGGGAAGAAGAGGGCAGGAG 1960
3' untranslated

L P S S F I A L A H G P R Q E N R F E E E G R R
GAGGGATGGGTCTCGGTGCTGAACCTTTGAGAACTTCCTACTAGGAATTGGAGGGGTGGAGTTTGAGA 2030
3' untranslated

R D G F S V L N L . E L P T R N W R G V S L R
ACTCCGTGCCCTTAACCTACATTTGCTGGCTCTCTTACGACTTAGGAGAAAAGATGATCTGGTCTTT 2100
3' untranslated

T P C P L T T F A G L L L R L R R K D D S G L
TCTTCAAGTTTGTTCACCTACAAACTCTTGGGCTTTCTGGGGAGGGATTGGAAGATATAAACAGACA 2170
3' untranslated

F F K F C F T Y K L L G F L G R D S E D I N P Q

FIG. 10 CONTINUED.

AACAAAAACAAACAAACCAACTACAGCAGTTCCAAGCTCGTTCTCACAAACACCTCTGAGACAGTCACAT 2240
3' untranslated
T K T N K P T T A V P S S F S Q T P L R Q S H

GTGGGCAATCTAAGGGAGGCAGGAAGCTCTACAGACTTTCTTGCAAACCTTCCCAGTTCTGTCGACAC 2310
3' untranslated
V G K S K G G R K L Y R L S C K P F P V L S T

TGCCAACACCTCCCCGCCAGAGACCTGGCCAGAGCCAAGAAAAGAGAAGCATGTGGTTTAAACAGAAAAA 2380
3' untranslated
L P T T S P P E T V P E P R K E K H V V . Q K N

CAAAACAAAACAAAACAAAATATATGTGTAAATCAACCTGTAGAAGGTAAAAACGGCAATGGAAAAGA 2450
3' untranslated
K T K Q N K K Y M C K S T C R R . K R Q V K R

TGAAGCTGGAAGGAGGGGCCAGTTGCCAAGATGGAACGAGAGCTGCCAGATCTTGCCCTCTGGATGACA 2520
3' untranslated
. S V K E G P S C Q D G T R A A R S C L L D D

AGAGGGGACATTGCAAGATGGCTGCCAGTCTAAAACGTCACCAGACCACAAGAGTAACATCACAGCCTTC 2590
3' untranslated
K R G H C K M A A S L K R H Q T T R V T S Q P S

GAAGAAAGGCCACAAGCTGTCTTCTTGCCCTCTAACTGAACATGCATGAAAAGTCAATAAACCCCTACTTT 2660
3' untranslated
K K G H K L S F C P L T E H A . K V N K P Y F

TTAATTTTTAAAAAAAAAAAAAAAAAAAAAAAAAATTCGCGGCCGC 2709
polyA tail + linker
L I F K K K K K K K K K F P R P

FIG. 11a.

AAGCTTGGCAGGAGGCTCGTGCCAAGCTGAGACCGTCATGCAGCTCCGAAATGAGTTAAGAGACAAGGA 70
LINKER ? open reading frame
A V H E A S C Q A E T V M Q L R N E L R D K E
GATGAAGCTGACAGATATCCGCTTAGAAGCTCTCAGTTCTGCCACCAGCTGGACCAGCTCCGGGAGGCC 140
open reading frame
M K L T D I R L E A L S S A H Q L D Q L R E A
ATGAACAGGATGCAGAGTGAAATAGAGAAGCTGAAAGCTGAGAATGATCGGCTGAAGTCAGAGTCTCAAG 210
open reading frame
M N R M Q S E I E K L K A E N D R L K S E S Q
GCAGTGGCTGCAGCCGGGCTCCTTCCCAAGTGTCATCTCTGCCTCCCGAGGCAGTCCATGGGCCCTCTC 280
open reading frame
G S G C S R A P S Q V S I S A S P R Q S M G L S
CCAGCACAGCTTGAACCTCACTGAGTCAACCAGCCTGGACATGTTGCTGGATGACACTGGTGAATGCTCG 350
open reading frame
Q H S L N L T E S T S L D M L L D D T G E C S
GCTCGGAAGGAAGGAGGCAGGCATGTTAAGATAGTTGTGAGCTTTTCAGGAGGAAATGAAGTGAAGGAGG 420
open reading frame
A R K E S G R H V K I V V S F Q E E M K V K E
ATTCAGACCACACCTCTTTCTTATTGGCTGCATTGGAGT TAGTGGAAGACGAAGTGGGATGTGCTCGA 490
open reading frame
D S R P H L F L I G C I G V S G K T K V D V L D
TGGGGTGGTTAGACGGCTGTTCAAAGAATACATCATTCATGTCGACCCAGTGAGTCAGCTAGGGCTGAAT 560
open reading frame
G V V R R L F K E Y I I H V D P V S Q L G L N
TCAGACAGCGTCTTGGCTACAGCATTGGAGAAATCAAGCGCAGCAACACTTCCGAAACACCGGAGCTGC 630
open reading frame
S D S V L G Y S I G E I K R S N T S E T P E L
TTCCTTGTTGGCTACTGCTTGGAGAGAACACGACCATCTCAGTGACTGTGAAAGGGCTCGCAGAAAACAG 700
open reading frame
L P C G Y L V G E N T T I S V T V K G L A E N S
CCTGGACTCAGTGCTGTTGAGTCCTTGATTCCCAAGCCATCCTGCAGCGCTACGTCCTCCTCCTGATA 770
open reading frame
L D S L V F E S L I P K P I L O R Y V S L L :
GAGCACCGTCGGATCATTCTCTCTGGCCCCAGCGGCACCTGGGAAAACCTACCTGGCCAACCGGCTGTC 840
open reading frame
E H R R : I L S G P S G T G K T Y L A N R L S

FIG. 11a CONTINUED.

AGTATATAGTGCTTCGAGAGGGACGGGAGTTGACAGACGGGGTTATCGCCACCTTTAACGTGGACCATAA 910
open reading frame
E Y I V L R E G R E L T D G V I A T F N V D H K
GTCCAGCAAGGAATTGCGCCAGTACCTGTCCAACCTTGCTGACCAGTGCAACAGTGAGAACAATGCTGTG 980
open reading frame
S S K E L R Q Y L S N L A D Q C N S E N N A V
GACATGCCCTCGTCATCATCCTGGACAACCTACACCACGTGAGCTCTCTGGGCGAGATCTTCAATGGGC 1050
open reading frame
D M P L V I I L D N L H H V S S L G E I F N G
TGCTCAACTGCAAGTACCACAAATGCCCTTACATAATTGGCACAATGAACCAGGCTACCTATCTCCCCCTT 1120
open reading frame
L L N C K Y H K C P Y I I G T M N Q A T Y L P F
TTATACTAATAATCTTATAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAATCTGCGGCCGC 1190
open reading frame LINKER-vector
Y T N N L I K K K K K K K K K K K K K K K F C G R

FIG. 11b.

| | | 10 | 20 | 30 | 40 | 50 |
|------------------------|--------------------------------------------------------|----|----|----|----|----|
| 1 1...50 UNC53_huma | AAGCTTGGCA CGAGGCCTCG TGGCAAGCTG AGACCGTCAT GCAGCTCCGA | | | | | |
| 5 1...48 | A W M E A S C Q A E T V M Q L R | | | | | |
| | 60 LINKER > 70 ORF (START) 80 90 100 | | | | | |
| 1 51...100 UNC53_huma | AATGAGTTAA GAGACAGGA GATGAAGCTG ACAGATATCC GCTTAGAAGC | | | | | |
| 5 49...98 | N E L R D K E M K L T D I R L E A | | | | | |
| | 110 120 130 140 150 | | | | | |
| 1 101...150 UNC53_huma | TCTCAGTCTT GCCCACCAGC TGGACCAGCT CCGGGAGGCC ATGAACAGGA | | | | | |
| 5 99...148 | L S S A H Q L D Q L R E A M N R | | | | | |
| | 160 170 180 190 200 | | | | | |
| 1 151...200 UNC53_huma | TGCAGAGTGA AATAGAGAAG CTGAAAGCTG AGAATGATCG GCTGAAGTCA | | | | | |
| 5 149...198 | M Q S E I E K L K A E N D R L K S | | | | | |
| | 210 220 230 240 250 | | | | | |
| 1 201...250 UNC53_huma | GAGTCTCAAG GCAGTGGCTG CAGCCGGGCT CTTTCCAAG TGTCCATCTC | | | | | |
| 5 199...248 | E S Q G S G C S R A P S Q V S I S | | | | | |
| | 260 270 280 290 300 | | | | | |
| 1 251...300 UNC53_huma | TGCCTCCCGG AGGCAGTCCA TGGGCTCTC CCAGCACAGC TTGAACCTCA | | | | | |
| 5 249...298 | A S P R Q S M G L S Q H S L N L | | | | | |
| | 310 320 330 340 350 | | | | | |
| 1 301...350 UNC53_huma | CTGAGTCAAC CAGCCTGGAC ATGTGCTGG ATGACACTGG TGAATGCTCG | | | | | |
| 5 299...348 | T E S T S L D M L L D D T G E C S | | | | | |
| | 360 370 380 390 400 | | | | | |
| 1 351...400 UNC53_huma | GCTCGGAAGG AAGGAGGCAG GCATGTTAAG ATAGTTGTCA GCTTCAGGA | | | | | |
| 5 349...398 | A R K E G G R H V K I V V S F Q E | | | | | |
| | 410 420 430 440 450 | | | | | |
| 1 401...450 UNC53_huma | GGAAATGAAG TGAAGGAGG ATTCCAGACC ACACCTCTTT CTTATTGGCT | | | | | |
| 5 399...448 | E M K W R E D S R P H L F L I G | | | | | |
| | 460 470 480 490 500 | | | | | |
| 1 451...500 UNC53_huma | GCATTGAGT TAGTGGCAAG ACGAAGTGGG ATGTGCTCGA TGGGGTGGTT | | | | | |
| 5 449...498 | C I G V S G K T K W D V L D G V V | | | | | |
| | 510 520 530 540 550 | | | | | |
| 1 501...550 UNC53_huma | AGAGGGCTGT TCAAAGAATA CATCATTCAT GTGAGCCAG TGAGTCAGCT | | | | | |
| 5 499...548 | R R L F K E Y I I H V D P V S Q L | | | | | |
| | 560 570 580 590 600 | | | | | |
| 1 551...600 UNC53_huma | AGGGCTGAAT TCAGACAGCG TTCTTGCTA CAGCATTGGA GAAATCAAGC | | | | | |
| 5 549...598 | G L N S D S V L G Y S I G E I R | | | | | |
| | 610 620 630 640 650 | | | | | |
| 1 601...650 UNC53_huma | GCAGGAACAG TTCCGAACA CCGAGCTGC TTCTTGTGG CTATCTGGTT | | | | | |
| 5 599...646 | R S H T S E T P E L L P C G Y L V | | | | | |
| | 660 670 680 690 700 | | | | | |
| 1 651...700 UNC53_huma | GGAGAGAACA CGACCATCTC AGTGAUTG AGAGGCTG CAGAAAACAG | | | | | |
| 5 649...696 | G E N T T I S V T V K G L A E N S | | | | | |

FIG. 11b CONTINUED.

| | | | | | |
|-----------------------------------------|---------------------------------------------------------|------|------|------|------|
| | 715 | 720 | 730 | 740 | 750 |
| 1 701..750 UNC53_huma 5 699..748 | CCTGGACTCA CTGGTGTTC AGTCCTTGAT TCCCAAGCCC ATCCTGCAGC | | | | |
| | L D S L V F E S L I P K P I L Q | | | | |
| | 765 | 770 | 780 | 790 | 800 |
| 1 751..800 UNC53_huma 5 749..798 | GCTACGCTCTC CCTCCTGATA GAGCACCCTC GATCATTTCT CTCTGGCCCC | | | | |
| | R Y V S L L I E H R R I I L S G P | | | | |
| | 810 | 820 | 830 | 840 | 850 |
| 1 801..850 UNC53_huma 5 799..848 | AGCGGCACTG GGAAAACCTA CCTGGCCAC CCGCTGTCTG AGTATATAGT | | | | |
| | S G T G R T Y L A N R L S E Y I V | | | | |
| | 860 | 870 | 880 | 890 | 900 |
| 1 851..900 UNC53_huma 5 849..898 | GCTTCGAGAG GGACGGGAGT TGACAGACGG GATTATCGCC ACCTTTAACC | | | | |
| | L R E G R E L T D G V I A T F N | | | | |
| | 910 | 920 | 930 | 940 | 950 |
| 1 901..950 UNC53_huma 5 899..948 | TGGACCATAA GTCCAGCAAG GAATTGGCC AGTACCTGTC CAACCTTGCT | | | | |
| | V D H K S S K E L R Q Y L S N L A | | | | |
| | 960 | 970 | 980 | 990 | 1000 |
| 1 951..1000 UNC53_huma 5 949..998 | GACCACTGCA ACAGTGAGAA CAATGCTGTG GACATGCCCC TCGTCATCAT | | | | |
| | D Q C N S E N N A V D M P L V I I | | | | |
| | 1010 | 1020 | 1030 | 1040 | 1050 |
| 1 1001..1050 UNC53_huma 5 999..1048 | CCTGGACAAC GTACACCAGG TGACCTCTCT GGGCGAGATC TTCAATGGCC | | | | |
| | L D N L H H V S S L G E I F N G | | | | |
| | 1060 PRIMER | 1070 | 1080 | 1090 | 1100 |
| 1 1051..1100 UNC53_huma 5 1049..1098 | TCTCAACTG GAAGTACCAC AAATGCCCTT ACATAATTGG CACAATGAAC | | | | |
| | L L N C K Y H K C P Y I I G T M N | | | | |
| | 1110 | 1120 | 1130 | 1140 | 1150 |
| 1 1101..1150 UNC53_huma 5 1099..1148 | CAGGCTACCT CTTCGACTCC CACCTGCCAG CTCACCATA ACTTCAGATC | | | | |
| | Q A T S S T P N L G L H H N F R W | | | | |
| | 1160 | 1170 | 1180 | 1190 | 1200 |
| 1 1151..1200 UNC53_huma 5 1149..1198 | GGTGCTTTGT GCCAACCACA CGGAGCCTGT GAAGGGTTTC CTGGCCGAT | | | | |
| | V L C A N H T E P V K G F L G R | | | | |
| | 1210 | 1220 | 1230 | 1240 | 1250 |
| 1 1201..1250 UNC53_huma 5 1199..1248 | TCTCGAGGAG GAAGCTCATG GAAACAGATA TCAGTGGGCG GGTGCGCAAT | | | | |
| | F L R R K L M E T E I S G R V R N | | | | |
| | 1260 | 1270 | 1280 | 1290 | 1300 |
| 1 1251..1300 UNC53_huma 5 1249..1298 | ATCGAGCTGG TAAAAATCAT TGACTGGATT CCAAGGTCT GGCATCACCT | | | | |
| | M I L V X I I D W I P K V W H H L | | | | |
| | 1310 | 1320 | 1330 | 1340 | 1350 |
| 1 1301..1350 UNC53_huma 5 1299..1348 | CAAGCGCTTC CTGAGGCTC ACAGTCTCTC GACGCTCAC ATCGCCCCC | | | | |
| | N R F L E A H S S S D V T I G P | | | | |
| | 1360 | 1370 | 1380 | 1390 | 1400 |
| 1 1351..1400 UNC53_huma 5 1349..1398 | GCTGCTTCCT GTCATGCCCC ATGATGTGCG ACGGCTCGAG AGTGTGGTTC | | | | |
| | R L F L S C P I D V D G S R V W F | | | | |

FIG. 11b CONTINUED.

| | | | | | | |
|-----------------------|-------------------------------------------------------|------|------|------|------|------|
| | | 1410 | 1420 | 1430 | 1440 | 1450 |
| 1 1401..1450 UNC53_hu | ACCGACTGT GGAACATATC CATTATCCCC TATCTCCTGG AAGCCGTCAG | | | | | |
| 5 1399..1448 | T D L W N Y S I I F Y L L E A V R | | | | | |
| | | 1460 | 1470 | 1480 | 1490 | 1500 |
| 1 1451..1500 UNC53_hu | AGAAGGACTC CAGCTCTATG GAAGCGCGCC CCCTGGGAG GATCCTGCCA | | | | | |
| 5 1449..1498 | E G L Q L Y G R R A P W E D P A | | | | | |
| | | 1510 | 1520 | 1530 | 1540 | 1550 |
| 1 1501..1550 UNC53_hu | AGTGGGTGAT GGACACATAT CCTGGGCAG CCAGCCACA ACAGCAGGAG | | | | | |
| 5 1499..1548 | K W V M D T Y P W A A S P Q Q H E | | | | | |
| | | 1560 | 1570 | 1580 | 1590 | 1600 |
| 1 1551..1600 UNC53_hu | TGGCTCCCC TCGTCAGTT ACGGCCTGAG GATGCGGCT TCGACGGCTA | | | | | |
| 5 1549..1598 | W P P L L Q L R P E D V G F D G Y | | | | | |
| | | 1610 | 1620 | 1630 | 1640 | 1650 |
| 1 1601..1650 UNC53_hu | CTCCATGGCT CGGAGGGAT CGACAAGCAA GCAGATCCCC CCGAGTATG | | | | | |
| 5 1599..1648 | S X P R E G S T S K Q M P P S D | | | | | |
| | | 1660 | 1670 | 1680 | 1690 | 1700 |
| 1 1651..1700 UNC53_hu | CTGAAGGTGA CCGCTGATG AACATGCTGA TGAGGCTGCA GGAGGCAGCC | | | | | |
| 5 1649..1698 | A E G D P L M N M L M R L Q E A A | | | | | |
| | | 1710 | 1720 | 1730 | 1740 | 1750 |
| 1 1701..1750 UNC53_hu | AACACTCCA GCGCCAGAG CTATGACAGC GACTCCACA GCACAGCCA | | | | | |
| 5 1699..1748 | N Y S S P Q S Y D S D S N S N S H | | | | | |
| | | 1760 | 1770 | 1780 | 1790 | 1800 |
| 1 1751..1800 UNC53_hu | TCAGATGAC ATCTGGACT CCTCTTTGGA GTGACTCTG TGCAGGGGC | | | | | |
| 5 1749..1790 | H D D I L D S S L E S T L | | | | | |
| | | 1810 | 1820 | 1830 | 1840 | 1850 |
| 1 1801..1850 UNC53_hu | CCGAGGCCA GCGCCCTCT CTCTCTCTCA CCGATTCCA CCGCATCCC | | | | | |
| 5 ---- | <== | | | | | |
| | | 1860 | 1870 | 1880 | 1890 | 1900 |
| 1 1851..1900 UNC53_hu | CACATCACC TGAAGATGAC TTCTGAGCC AGCCCCAGCC ACAGCCTTAG | | | | | |
| 5 ---- | <== | | | | | |
| | | 1910 | 1920 | 1930 | 1940 | 1950 |
| 1 1901..1941 UNC53_hu | AGCTCGGGA ACACCGAGAC CCCCCTCTT CAGCCTGAC | | | | | |
| 5 ---- | <== | | | | | |

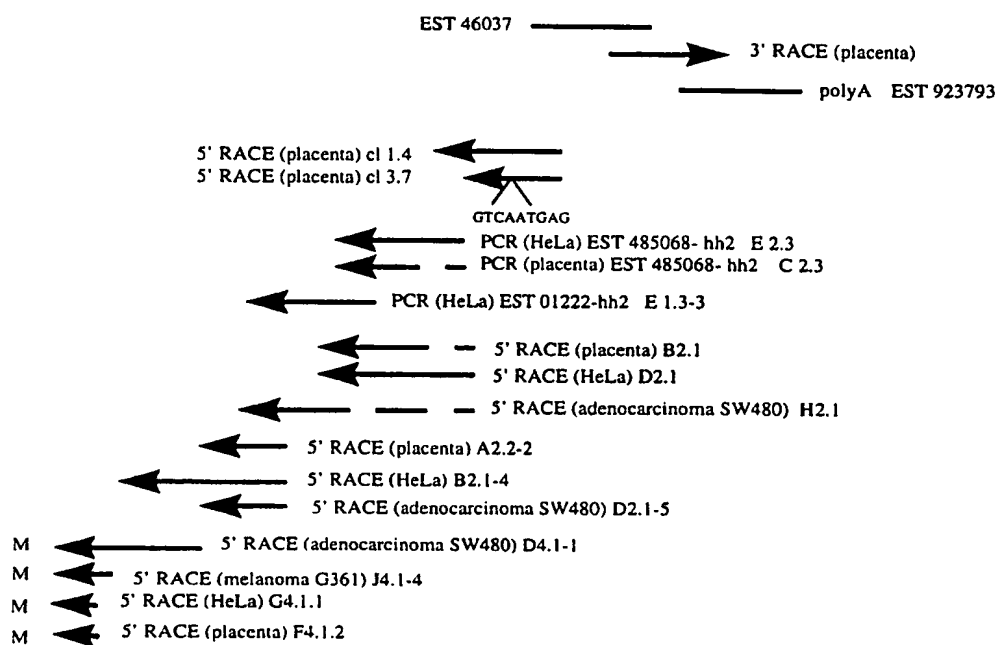


Figure 11c

11 K E E P K E D P S G A A V P E M P K K S S K I A S F
1281 CAGAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
GAGAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 P K G G K L N S A K K E P M A P S H S G I P K P G M
1361 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 K S N P O R S P S A P A P S K E G E R S R S G K L S
1441 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 G L P Q Q K P Q L D G R H S S S S S L A S S E G K
1521 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 G P G G T T L N H S I S S Q T V S G S V G T T Q T T G
1601 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 S N T V S V Q L P Q P Q Q Q Y N H P N T A T V A P F
1681 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 Y R S Q P Q T E G N V P A E S S T G V S V E P S H
1761 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 F T K T G O P A L E E L T G E D P E A R R L R T V X N
1841 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 I A D L R Q N L E E T M S S L R G T Q V T H S T L E
1921 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 T F D T N V T E H S G R S I L S L T G R P T P L S
2001 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 M R L G Q S S P R L O A G D A P S H G N G Y P P R A H
2081 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 A S R F I N T E S G R Y V Y S A P L R R O L A S R G
2161 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 S V C H V D V S D K A G D E M D L E G I S M D A P G
2241 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 Y M S D G O V L E K N I R T D I T S G Y H T D G G L
2321 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 G L Y T R E L N R L P D G H A V V R E T L Q R N T S
2401 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 G L G D A D S M D D S S S V S S G I S D T I D N L S
2481 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG

FIGURE 11d

11 I Y T D W A H N Y L T K S G H K R L I K D L O O D V
1 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 D G V L L A O I L O V V A N E K I E D I N G C P K N
61 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 P S O N I E H I D A C L H P L A A K C I N I O G L S A
141 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 E E I R N G H L K A I L G L F S L S R Y K O O Q
211 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 P O K Q H L S S P L P A V S Q V A G A P S Q C O A
321 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 G T P O Q V P T P Q A P C O P H O P A P H O Q S K
401 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 A O A E M O S R L S G P T A R V S A A G S E A K T R
481 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 G S T A M H R R S Q S F H N Y D K S E P V T S P P
561 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 P P G S H E K E P L A S A S H P C N S D N A P A
641 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 S L E S C S S T P T N C S T S S A I P O P G A A T
721 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 P M R S K S L S V K H S A T V S H L S V K P P G P E
801 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 A P P T P E A H K P A P N H O K S H L E K L K L F N
881 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 S K G S E A G E G P G S R D T S C E R L E T L P S
961 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 E S E E L E A A S R M L T T V G P A S S S P K I A
1041 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 L E G I A O R T F S R A L T N K E S S L K G N E K E K
1121 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
11 E K O R E K D K E K S K D L A K R A S V T E R L D
1201 ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG
ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG ACCAGAG

[illegible][illegible]

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Fig 11c

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Page 1

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+1 G T R V T I G P R L F L S C P I D V D G S R V W F T D
1 GGCACGAGGG TTACCATCGG CCCCCGGGCTC TTCCTGTCAT GCCCCATCGA TGTGGACGGC TCGAGAGTGT GGTTCACCGA
CCGTCTCCC AATGGTAGCC GGGGGCCGAG AAGGACAGTA CGGGGTAGCT ACACCTGCCG AGCTCTCACA CCAAGTGGCT
.....
+1 L W N Y S I I P Y L L E A V R E G L Q L Y G R R A P
81 CTTGTGGAAC TATTCATTA TCCCCTATCT CCTGGAAGCC GTCAGAGAAG GACTCCAGCT CTATGGAAGG CGCGCCCCCT
GAACACCTTG ATAAGGTAAT AGGGGATAGA GGACCTTCGG CAGTCTCTTC CTGAGGTGCA GATACCTTCC GCGCGGGGGA
.....
+1 W E D P A K W V M D T Y P W A A S P Q Q H E W P P L L
NcoI
-----
161 GGGAGGATCC TGCCAAGTGG GTGATGGACA CATATCCATG GGCAGCCAGC CCACAACAGC ACGAGTGGCC TCCCCTGCTG
CCCTCCTAGG ACGGTTTCACC CACTACCTGT GTATAGGTAC CCGTCGGTCG GGTGTTGTCT TGCTCACCAG AGGGGACGAC
.....
+1 Q L R P E D V G F D G Y S M P R E G S T S K Q M P P S
241 CAGTTACGGC CTGAGGATGT CGGCTTCGAC GGCTACTCCA TGCCCTCGGA GGGATCGACA AGCAAGCAGA TGCCCCCAG
GTCAATGCCG GACTCCTACA GCCGAAGCTG CCGATGAGGT ACGGAGCCCT CCCTAGCTGT TCGTTCGTCT ACGGGGGGTC
.....
+1 D A E G D P L M N M L M R L Q E A A N Y S S P Q S Y
321 TGATGCTGAA GGTGACCCGC TGATGAACAT GGTGATGAGG CTGCAGGAGG CAGCCAACTA CTCCAGCCCC CAGAGCTATG
ACTACGACTT CCACTGGGCG ACTACTTGTA CGACTACTCC GACGTCCTCC GTCGGTTGAT GAGGTCGGGG GTCTCGATAC
.....
+1 D S D S N S N S H H E D I L D S S L E S T L * Q G P G
401 ACAGCGACTC CAACAGCAAC AGCCATCAGC AAGACATCTT GGACTCCTCT TTGGAGTCCA CTCTGTGACA GGGGCCCCGA
TGTCGCTGAG GTTGTCGTTG TCGGTAGTGC TTCTGTAGAA CCTGAGGAGA AACCTCAGGT GAGACACTGT CCCCAGGCGT
.....
+1 A Q R P P L L L T A F H L H P P H H P E D D F L S Q P
481 GCCCAGCGCC CTCCTCTTCT CCTCACCGCA TTCCACCTGC ATCCCCCACA TCACCCTGAA GATGACTTCC TGAGCCAGCC
CGGGTCGCGG GAGGAGAAGA GGAGTGGCGT AAGGTGACG TAGGGGGTGT AGTGGGACTT CTACTGAAGG ACTCGGTCCG
.....
+1 P A T A L E L R E H R D P P S F S L D L G A G I P G
561 CCCAGCCACA GCCTTAGAGC TCGGGGAACA CCGAGACCCC CCGTCCTTCA GCCTCGACCT GGTGTCAGGC ATCCCGGGCC
GGGTCTGGTGT CGGAATCTCG ACGCCCTTGT GGCTCTGGGG GGCAGGAAGT CGGAGCTGGA CCCACGTCCG TAGGGCCCGG
.....
+1 Q L P A D R F L P Q R E L H Y L L L Y F N Y C F A L L
641 AGCTGCCTGC GGACCGCTTC CTTCCACAGC GAGAAGTCCA CTACCTTCTG TTGTACTTTA ATTATTGTTT TGCCTTGTTG
TCGACGGACG CCTGGCGAAG GAAGGTGTCG CTCTTGACGT GATGGAAGAC AACATGAAAT TAATAACAAA ACGGAACAAC
.....
+1 L * P P * D T E D T S R E R I I A V E M K K K K K K K
721 CTGTGACCTC CTAAGACAC TGAAGATACT TCTCGGGAAG GGATCATCGC CGTTGAAATG AAAAAAAAAA AAAAAAAAAA
GAACTGGAG GGATTCTGTG ACTTCTATGA AGAGCCCTTT CCTAGTAGCG GCAACTTTAC TTTTTTTTTT TTTTTTTTTT
.....
+1 K K K K K N E G G R K L
801 AAAAAAAAAA AAAAAAACG AAGGCGGCGG CAAGCTT
TTTTTTTTTT TTTTTTTTGC TTCCGCGGCG GTTCGAA
```

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hh2UNC53_5'Race_cl1.4_CON_REV.seq

Page 1

+1 I P L T I G L E R P P R Q V P H L D R N T L P K K G L
1 ATCCCACTCA CTATAGGGCT CGAGCGGCGG CCCAGGCAGG TCCCGCACCT TGATAGGAAC ACTTTGCCTA AGAAAAGGACT
TAGGGTGAGT GATATCCCGA GCTCGCCGCG GGGTCCGTCC AGGGCGTGGA ACTATCCTTG TGAAACGGAT TCTTTCTCTGA

+1 R Y T P T S Q L R T Q E D A K E W L R S H S A G G L
81 CAGGTATACT CCCACCTCCC AGCTTCGCAC GCAAGAAGAT GCAAAAGAAT GGTTACGGTC CCATTCTGCA GGAGGCCTTC
GTCCATATGA GGGTGGAGGG TCGAAGCGTG CGTTCTTCTA CGTTTTCTTA CCAATGCCAG GGTAAGACGT CCTCCGGAAG

+1 Q D T A A N S P F S S G S S V T S P S G T R F N F S Q
161 AGGACACCGC TGCCAATTCC CCCTTTTCTT CTGGCTCCAG CGTGACTTCT CCCTCCGGA CAAGATTCAA CTTTTCCTCAG
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+1 L A S P T T V T Q M S L S N P T M L R T H S L S N A D
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+1 S T P E E K C Q S E I R K L R R E L D A S Q E K V S A
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+1 L T T Q L T A N A H L V A A P E Q S L G N M T I R L
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+1 N A A A Q A A I N G V I N T P E L N C K G N G T A Q S
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+1 G S N I E S D S K K K K R K N W L R S S F K Q A F G K
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[illegible]

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 13 241 G R Y V S A P L R O L A S R G S S V
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1 L G F Q S V S S L H T S C E S I D I S L S S G G V P
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 5 1361 S H N S S T G L I A S S R D D S L T P F V R T N S V R
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25ATVSMLSVKPPGPPEAPRPTPEANMKPA

.....2 P N N Q K S M L E K L K L P N S' K O G S K A G E G P

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ТАБАЧНИКОВ СЕРГЕЙ АЛЕКСАНДРОВИЧ ТРОФИМЕНКО АЛЕКСАНДР ГИРИЧЕНКО МАТЕВЦЕВ

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1 GCACGACCT TCCACGAG CECTCTTGA CCGACGACCT GACTCTTAC GACGACCTA MGACGACC CATTCACCA

2 A V P E M P K K S S K I A S P I P K G C K L N S A R K
6 C T T T A C C C G A G A T T C C A A A A G T C T C A G T T T C C A A G G G G G G T A C T T A C A G T T C C A A

2 E P M A P S H S G I P K P G M K S M P G K S P S A P

2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65 66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 92 93 94 95 96 97 98 99 100 101 102 103 104 105 106 107 108 109 110 111 112 113 114 115 116 117 118 119 120 121 122 123 124 125 126 127 128 129 130 131 132 133 134 135 136 137 138 139 140 141 142 143 144 145 146 147 148 149 150 151 152 153 154 155 156 157 158 159 160 161 162 163 164 165 166 167 168 169 170 171 172 173 174 175 176 177 178 179 180 181 182 183 184 185 186 187 188 189 190 191 192 193 194 195 196 197 198 199 200 201 202 203 204 205 206 207 208 209 210 211 212 213 214 215 216 217 218 219 220 221 222 223 224 225 226 227 228 229 230 231 232 233 234 235 236 237 238 239 240 241 242 243 244 245 246 247 248 249 250 251 252 253 254 255 256 257 258 259 260 261 262 263 264 265 266 267 268 269 270 271 272 273 274 275 276 277 278 279 280 281 282 283 284 285 286 287 288 289 290 291 292 293 294 295 296 297 298 299 300 301 302 303 304 305 306 307 308 309 310 311 312 313 314 315 316 317 318 319 320 321 322 323 324 325 326 327 328 329 330 331 332 333 334 335 336 337 338 339 340 341 342 343 344 345 346 347 348 349 350 351 352 353 354 355 356 357 358 359 360 361 362 363 364 365 366 367 368 369 370 371 372 373 374 375 376 377 378 379 380 381 382 383 384 385 386 387 388 389 390 391 392 393 394 395 396 397 398 399 400 401 402 403 404 405 406 407 408 409 410 411 412 413 414 415 416 417 418 419 420 421 422 423 424 425 426 427 428 429 430 431 432 433 434 435 436 437 438 439 440 441 442 443 444 445 446 447 448 449 450 451 452 453 454 455 456 457 458 459 460 461 462 463 464 465 466 467 468 469 470 471 472 473 474 475 476 477 478 479 480 481 482 483 484 485 486 487 488 489 490 491 492 493 494 495 496 497 498 499 500 501 502 503 504 505 506 507 508 509 510 511 512 513 514 515 516 517 518 519 520 521 522 523 524 525 526 527 528 529 530 531 532 533 534 535 536 537 538 539 540 541 542 543 544 545 546 547 548 549 550 551 552 553 554 555 556 557 558 559 560 561 562 563 564 565 566 567 568 569 570 571 572 573 574 575 576 577 578 579 580 581 582 583 584 585 586 587 588 589 590 591 592 593 594 595 596 597 598 599 600 601 602 603 604 605 606 607 608 609 610 611 612 613 614 615 616 617 618 619 620 621 622 623 624 625 626 627 628 629 630 631 632 633 634 635 636 637 638 639 640 641 642 643 644 645 646 647 648 649 650 651 652 653 654 655 656 657 658 659 660 661 662 663 664 665 666 667 668 669 670 671 672 673 674 675 676 677 678 679 680 681 682 683 684 685 686 687 688 689 690 691 692 693 694 695 696 697 698 699 700 701 702 703 704 705 706 707 708 709 710 711 712 713 714 715 716 717 718 719 720 721 722 723 724 725 726 727 728 729 730 731 732 733 734 735 736 737 738 739 740 741 742 743 744 745 746 747 748 749 750 751 752 753 754 755 756 757 758 759 760 761 762 763 764 765 766 767 768 769 770 771 772 773 774 775 776 777 778 779 780 781 782 783 784 785 786 787 788 789 790 791 792 793 794 795 796 797 798 799 800 801 802 803 804 805 806 807 808 809 810 811 812 813 814 815 816 817 818 819 820 821 822 823 824 825 826 827 828 829 830 831 832 833 834 835 836 837 838 839 840 841 842 843 844 845 846 847 848 849 850 851 852 853 854 855 856 857 858 859 860 861 862 863 864 865 866 867 868 869 870 871 872 873 874 875 876 877 878 879 880 881 882 883 884 885 886 887 888 889 890 891 892 893 894 895 896 897 898 899 900 901 902 903 904 905 906 907 908 909 910 911 912 913 914 915 916 917 918 919 920 921 922 923 924 925 926 927 928 929 930 931 932 933 934 935 936 937 938 939 940 941 942 943 944 945 946 947 948 949 950 951 952 953 954 955 956 957 958 959 960 961 962 963 964 965 966 967 968 969 970 971 972 973 974 975 976 977 978 979 980 981 982 983 984 985 986 987 988 989 990 991 992 993 994 995 996 997 998 999 1000 1001 1002 1003 1004 1005 1006 1007 1008 1009 1010 1011 1012 1013 1014 1015 1016 1017 1018 1019 1020 1021 1022 1023 1024 1025 1026 1027 1028 1029 1030 1031 1032 1033 1034 1035 1036 1037 1038 1039 1040 104

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11/24/97 03:58PM B2.1-4 (hh2UNC53 5:PACE 02/04/97) Page 1

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[illegible]

B2.1-4 (hh20nc53 5 RACE 02/08/97) Page 2

2104/971

Page 2

2001 TAAATTTTAAACCAAGCAATTT C
ATTATTAATTATTTTATTTTAAAG

ESCOR I

2001 TCACTTCA | AGCGTAAT C
ACTTAAATTT TTGAG TTAA G

References

*) S L * G S S G R P G R C S S T P T H C

1 GATITICA GATITICIT TATITICIT AGGATITICAG GAGGATITICAG GAGGATITICAG TATITICIT
GATITICIT GATITICITAG ATATITICITAG TATITICITAG GAGGATITICAG GAGGATITICAG ATATITICITAG

*) S T S A I P O P G A A T R P M R S K S L S V K H S A
81 AGGATITICIT GAGGATITICIT GAGGATITICIT GAGGATITICIT GAGGATITICIT GAGGATITICIT
TATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) T V S M L S V K P P G P E A P M P T P E A N K P A P
161 GAGGATITICIT AGGATITICIT TATITICITAG TATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG TATITICITAG AGGATITICITAG AGGATITICITAG AGGATITICITAG AGGATITICITAG

*) H R O K S M L E K L K L F N S K G G S K A G E G P O S
241 AGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
TATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) R D T S C E R L E T L P S P E E S E L E A A S R M L
311 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) T T V G P A S S S P K I A L K G I A Q R T P S R A L
401 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) T N K K S S L K G N E K E K E K O O R E K D K E K S K
481 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) O L A K R A S V T E R L D L K E E P R E D P S G A A V
561 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) P E M P K R S S K I A S P I P K G G L N S A K E
641 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) P H A P S H S G I P K P G M R S M P G K S P S A P A P
721 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) S K E G E R S R S G K L S S G L P O Q X P O L D O R H
801 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) S S S S S L A S S E G K G P O G T T L N H S I S S
881 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) Q T V S G S V G T T O T T G S N T V S V O L P O P Q Q
961 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) Q V H P M T A T V A P P L V R S O T D T E G N V T A
1041 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) E S S S T G V S V E P S H P T K T G O P A L E E L T
1121 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) G E D P E A R R L R T V K H I A O L R Q N L S E T V S
1201 GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG
GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG GAGGATITICITAG

*) S L R G T O V T K P H
1281 AGGATITICITAG GAGGATITICITAG TATITICITAG
TATITICITAG GAGGATITICITAG TATITICITAG TATITICITAG

[illegible]

Fig. 11f.

1 CTGACAGU TGTTCAGCA AGGTTTAT CTTCAGCC GAGACAGC CTCTTTCG AGTTTACCA GCTTCAGT
GAGTTTCT AGACTCTT TCCAMAH CAAGTCCG GCTCTTCC GACAAACG TCAAGCTG CGAGGCTAA

81 GCGCGGCA AGCAGGCA TCGCTGCA GAGACAGC ATCGCTCT TCTCTTCC GCTTCCCA GCGACAGC
GCGGCTCT TGTCTCTC AGCGACT CTCTTTCG TACAGCCA AGCAGGCT GCGAGGCT CTCTCTCC

83 K O T L F A E T S T A F I I T K P L P C S D N E F O

161 CAGACACA CTCTTCTG AGCTTCAC AGCTTCTT ATACAMAC CTCTCTCC GCTACCAT GAGTTCCG
GCTCTCTG GCGAGGAC TGTAGAGC TGTAGAGC TATCTTTC GAGAGCTC GAGCTCTTA CTGAGCTG

241 Q H A L P R A S V E H L E * Q L D F I E C L P C T

241 AAATTCG AGCTCTCC AGAGCTCT TACAGCTC TACAGCA CTGAGCTTA TTAGCTCTT ACATCCAC
TTATCTCC TATAGAGC TCTCTGAC ATCTCTCA TCTTCTT GACTTCAAT ACTACAAA TATAGCTG

321 K P W A K H F I C R L F V F T A H P V G R Y N Y P H S

321 AGGCTCCG CTAAAGCT CATCTCAG CTCTCTCT TACGCGAA CCGATAGCT AGGTATCT ATCCGACT
TTCGAGCC GTAGCTCC GAGAGCTCC GAGAGGAG ATCTCTTT GCGTATCA TCGATTTCA TACGCTAG

401 A D A E T E A Q S V L V A K Q A H Q E A R K W P H L

401 TCAAGTCA GAGAGGAG CAGAGGCT TTTCTAGT AAAAGCTC ACTAGAGCC TACAGGCT GAGAGCTAG
AGCTTCTT CTCTCTCC GAGAGCTCA TTTCTAG TTTCTAG TCTCTCCG ATCTCTCC GCTCTGATC

461 A G P P D S T N C L P L L C C H Q E C D S K F F L P S

461 CTGCTCTCC TACTCTCC ACTCTCTCC CTCTCTCT TCGATCGAA GATCTGCT CAGCTTTT CTCTCTCT
GAGCGGGG ACTAGCTG TTAGCGAG TTAGCGAG GAGAGGAG AGCTTCTT CTACTCTA GCTTCAAAA GAGAGGAG

561 G S H S G P T L L S H Q I Y T D W A N H Y L A K S G H

561 GATCTACT CTGCTTAC TCTCTTCC AGAGGCTC AGAGGCTC GCGATCAT TACTAGCA ATCTCTCA
CTTAGCTCA GAGGAGCT AGAGGCTG TGTCTTCA TGTCTTCC GCGTATTA ATGATCTT TTAGCTCT

641 K R L I K D L Q O E S R I P A H W R P L L V D P S S

641 CAGCTCTC ATAGAGCT TCGAGGAA AGCGGAT CAGGACT GCGGCTCT ACTAGCTG CAGAGCTG
CTTCCAGAG TACTCTCT AGCTCTCT TCGCTTAA GCTCTCTA CCGCGGCA TACTACTA GCTCTGAC

721 V P K L G V I M V I V

721 TACGAGCT TCGGATCT ATCTCTAG TGT

ATCTCTCA ATCTCTAG TACTCTATC ACA

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1 RRRAGAGCA GGGGGTAA AAGGTTTTH AUAAGAGG GATCCCTTG NGGGGTHT TTTTNGCC CCGKGGTTG
AGCTTCTCT CCGGATTT TTRGAAAT TTTTTRCC CTAGAGAC ACCCGAAA AAAGAGGG GGGKCGAAC
.....
81 TAAAGGGG TCCCTTAGG ATTTTHAG GAGGGGAG GAGGGGAG ACCTTTCT TGGGGGGH GTCAAGGCG
ATTTTCCC AGGAGTAC TAAAGATTC CTTCCTTG CTCCCTTG TGAAGAGA ACCCGCGH CACTTTGCG
.....
161 HCAACAGCT GGGGGGAG GCGAGGAG TTTTCTTT AUAAGGGA AATTGGAG AAGAGGCG TTTGGTAAAG
HTGGTGGG CCGGCTCG KCGTTGTH AUAAGGAA TTTTCTCT TAAAGCTC TTTTCCGC AAACGATTC
.....
241 TGGGGAGT TTTGGGNC ATNAGATG AGAGTAGG TGGATTCG CTGAGGCG GCGAATGC INGGTCTCT
HACCTGCA AAGCGGCG TAAATGTA TGTGATAC AGCTAGCG GATTTGCG CCGTTACG INCCGAGGA
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321 AGCTGCGG TTTTTCGC GAGGTGGA AGGAGAGG TTAAGGAT TCGATTAH CTGGTTAA TGAAGACTT
TTAAAGCG AAGAGAGG CTGAGGCT TCGGTTCC AATTTCTH AGGTAATT GAGCGAATT ACTTTGAA
.....
401 TGGTTTTT HTATTAHA ATATTAHA TAAAGGAT TTTCTTCC CAGAAAAA AAGAGAGA AAGTAAACA
ACTAAAAA HTATTAAT TATTAATT ATTTTTHA AAGGAGGG GTTTTTTT TTTTCTCT TTTATTTG
.....
481 HAAATATA HAAAGAGH GAAAGAGH AATCAAAA NAAAGAGG GCGTTGAT TTTAGCTC TAAAGGATG
HTATTAAT HTTTTTHG CTTTTCTH TTTAGTTT TTTTTCGC CCGAATTA AATTTGAGG ATCTGTAC
.....
561 V R P H S L S R D R V V R V P R A P I V S . Y F G V
TGTGCGCT CAGAGCTT CCGGAGAG AGTGTTCG GTCCCGGAG CCGCATCT GTCTTATC TTTGGGTTG
ACAGCGGA GTTGGGCA GGGCTGTC TCAGAGCG GAGGGGCT GGGGTACA CAGACTATG AAACCGAG
.....
641 H M A I D L Y C G L A C L W G K H E P R I Y T D W A N
AGATGCTAT AGATTTTAC TGGTTTGG CTGTCTTG GAGAGGAT GAGCGCGA TCTACAGA CTGGGCAAT
TGTAGGATA TTAAGATG ACCGAGCC GAGAGAGC CCGTTTCTA CTGGGCTT AGATGCTC GAGCGTTA
.....
721 H Y L A K S G H K R I I K D F Q Q E S R I L Q I S I T
CATTAAGT CGAATCGG CGAGAGCT ATATGAGG ATTTGAGA AAGAGGCA ATTTGAGA TATCATAC
GTATGATC GTTTAGCC GTTTTCCA TATGATTC TAAAGTCT TTTTCTCT TAAAGCTC ATAGTAGTG
.....
801 X A A A B A C X . R P I R P I V S C I Q
ATGCGGCT GTGAGAGT GAGTACAG GCGATTCG CCAATAGCA GTTTATACA AT
TAAAGCGG GAGTACGA CTAATATC CCGTAGAG GATATCAT CAGATATG TA
.....

Figure 12. Tblastn search of the EST division of Genbank with 680aa starting at the c-terminus of the alfa-actinin domain of Hu-UNC-53/2.

LOCUS AA418158 610 bp mRNA EST 19-MAY-1997
DEFINITION zv97dl2.r1 Soares NhBMPu S1 Homo sapiens cDNA clone 767735 5'.
ACCESSION AA418158
NID g2079968
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
 Eukaryotae; mitochondrial eukaryotes; Metazoa; Chordata;
 Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae;
 Homo.
REFERENCE 1 (bases 1 to 610)
AUTHORS Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,
 Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J.,
 Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theising, B.,
 White, Y., Wylie, T., Waterston, R. and Wilson, R.
TITLE WashU-Merck EST Project 1997
JOURNAL Unpublished (1997)
COMMENT
 Contact: Wilson RK
 WashU-Merck EST Project
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 This clone is available royalty-free through LLNL ; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 Seq primer: -28ml3 rev2 ET from Amersham
 High quality sequence stop: 492.
FEATURES Location/Qualifiers
source 1..610
 /organism="Homo sapiens"
 /note="Organ: mixed (see below); Vector: pT7T3D-Pac
 (Pharmacia) with a modified polylinker; Site_1: Not I;
 Site_2: Eco RI; Equal amounts of plasmid DNA from three
 normalized libraries (melanocyte 2NbEM, pregnant uterus
 NbHPU, and fetal heart NbHH19W) were mixed, and ss circles
 were made in vitro. Following HAP purification, this DNA
 was used as tracer in a subtractive hybridization
 reaction. The driver was PCR-amplified cDNAs from pools of
 5,000 clones made from the same 3 libraries. The pools
 consisted of I.M.A.G.E. clones 260232-265223,
 340488-345479, and 484488-489479."
 /clone="767735"
 /clone_lib="Soares NhBMPu S1"
 /tissue_type="Pooled human melanocyte, fetal heart, and
 pregnant uterus"
 /lab_host="DH10B"
mRNA <1..>610

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                                /clone="5D16"
                                /clone_lib="Zebrafish ICRFzf1s"
                                /sex="mixed"
                                /tissue_type="pooled 26-somite embryos"
                                /lab_host="XL1-blue MRF"
                                complement(<1..>418)
mRNA
BASE COUNT      108 a      87 c      78 g      145 t
ORIGIN
    1 tttacatttt ttgaggaaga tgctaattggt ctattctgat tcaatgattt atgctaagct
   61 aagctaaaaat gctcctgtca aatcctgaga tcagctgaat gaattaaaaa ttgggtaaaa
  121 ctcaactgtc taactctagg ggagttgtaa aatgggccta tttccctaaa aagtaatgtt
  181 actttaagag catgatggtc caccagtttc actgtctaaa ttttgttatt ccataagcta
  241 atcttctctg ggcattttga cgattttaac actaacctgt gggtaatctg cgtcccccgt
  301 aaactggaca tggtttcttc cagattctgt ctcagatcag caatgttctt cactgtacgc
  361 atccgtctag tttctggatc ttctctgag atctcctcca ggcactgttt ggcgggtct
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gb|AA495042|AA495042 fa05f06.s1 Zebrafish ICRFzf1s Danio rerio cDNA
      clone 5D16 3'
      Length = 418

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Minus Strand HSPs:

Score = 195 (87.9 bits), Expect = 9.9e-18, P = 9.9e-18
 Identities = 37/46 (80%), Positives = 42/46 (91%), Frame = -3

```

Query:   627 TGQPALEELTGEDPEARRLRITVKNIADLRQNLLEETMSSLRGTVQVTH 672
          T + LEE++GEDPE RR+RTVKNIADLRQNLLEETMSSLRGTVQ+TH
Sbjct:   416 TAKQCLEEISGEDPETRRMRTVKNIADLRQNLLEETMSSLRGTVQITH 279

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MOUSE 2

```

LOCUS      AA208994      527 bp      mRNA      EST      18-FEB-1997
DEFINITION mw75e12.r1 Soares mouse NML Mus musculus cDNA clone 676558 5'.
ACCESSION  AA208994
NID        gl807004
KEYWORDS   EST.
SOURCE     house mouse.
  ORGANISM Mus musculus
            Eukaryotae; mitochondrial eukaryotes; Metazoa; Chordata;
            Vertebrata; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
            Mus.
REFERENCE  1 (bases 1 to 527)
  AUTHORS  Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,
            Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,
            Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,
            Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and
            Waterston,R.
  TITLE    The WashU-HHMI Mouse EST Project
  JOURNAL  Unpublished (1996)
COMMENT
  Contact: Marra M/Mouse EST Project
  WashU-HHMI Mouse EST Project
  Washington University School of MedicineP
  4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

```

Tel: 314 286 1800

Fax: 314 286 1810

Email: mouseest@watson.wustl.edu

This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:416262

Putative full length read

vector to vector length is 535

Seq primer: -28ml3 rev2 ET from Amersham

High quality sequence stop: 478.

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FEATURES
    source                Location/Qualifiers
                        1..527
                        /organism="Mus musculus"
                        /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                        polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
                        was primed with a Not I - oligo(dT) primer [5'
                        TGTACCAATCTGAAGTGGGAGCGGCCGCGAATCTTTTTTTTTTTTTTTT 3'];
                        double-stranded cDNA was ligated to Eco RI adaptors
                        (Pharmacia), digested with Not I and cloned into the Not I
                        and Eco RI sites of the modified pT7T3 vector. Library
                        constructed and normalized by Bento Soares and M.Fatima
                        Bonaldo."
                        /clone="676558"
                        /clone_lib="Soares mouse NML"
                        /tissue_type="Liver"
                        /lab_host="DB10B"
    mRNA                 <1..>527
BASE COUNT             151 a    139 c    136 g    101 t
ORIGIN
    1 tgtctctgga tgagaagagc cgaacaatga gtcggtcagg ctccttccgg gatggggttg
   61 aggaagttca tggatcctcc ctgtccttgg tttccagcac atcctccatc tactccacgc
  121 cagaagaaaa atgccagtca gagattcgaa agctgaggcg agacgtggat gcctcccagg
  181 aaaaggtgtc tgcgctgact acccagctga ctgcaaatgc tcaccttgtg gcagccttgc
  241 agcagagtct gggaaacatg accatcaggc tacagagttt aactatgacc gctgagcaga
  301 aggattcaga actgaacgag ttaagaaaaa ccacgcagct gctgaagaaa cagaatgcag
  361 ctgcccaggc tgccattaat ggagtgatta acacgccaga gctcaactgc aaaggaaatg
  421 gcagtgccag gctacagacc tacgcattcc cagcaacact cctccgacag tgtctccagt
  481 atcaatagcg ccaccagcca ctcaagtgtg ggcagcaaca tagagag

```

gb|AA208994|AA208994 mw75e12.r1 Soares mouse NML Mus musculus cDNA
clone 676558 5'
Length = 527

Plus Strand HSPs:

Score = 541 (243.9 bits), Expect = 2.3e-76, Sum P(2) = 2.3e-76
Identities = 110/143 (76%), Positives = 114/143 (79%), Frame = +3

Query: 1511 SLDEKSRMSRSGSFRDGFEEVHGXXXXXXXXXXXXXXXXXPEEKQSEIRKLRLRELDASQE 1570
SLDEKSRMSRSGSFRDGFEEVHG PEEKQSEIRKLRR++DASQE
Sbjct: 3 SLDEKSRMSRSGSFRDGFEEVHGSSLSVSTSSSIYSTPEEKQSEIRKLRRDVDASQE 182
Query: 1571 KVSALTTLTANAHLVAAFEQSLGNMTIRLQSLTMTAEQKDELNELRKTIEXXXXXXXXX 1630

Sbjct: 183 KVSALTTQLTANAHLVAAFEQSLGNMTIRLQSLTMTAEQKDSELNELRKTI
 362 KVSALTTQLTANAHLVAAFEQSLGNMTIRLQSLTMTAEQKDSELNELRKTIELLKKQNAA

Query: 1631 XXXXXXGVINTPELNCKGNGTAQ 1653
 GVINTPELNCKGNG+A+

Sbjct: 363 AQAAINGVINTPELNCKGNGSAR 431

Score = 116 (52.3 bits), Expect = 2.3e-76, Sum P(2) = 2.3e-76
 Identities = 24/25 (96%), Positives = 25/25 (100%), Frame = +1

Query: 1661 QHSSDSVSSINSATSHSSVGSNIE 1685
 +QHSSDSVSSINSATSHSSVGSNIE

Sbjct: 451 QHSSDSVSSINSATSHSSVGSNIE 525

LOCUS AA049124 337 bp mRNA EST 09-SEP-1996
 DEFINITION mj46f04.r1 Soares mouse embryo NbME13.5 14.5 Mus musculus cDNA
 clone 479167 5'.
 ACCESSION AA049124
 NID gl528794
 KEYWORDS EST.
 SOURCE house mouse.
 ORGANISM Mus musculus
 Eukaryotae; mitochondrial eukaryotes; Metazoa; Chordata;
 Vertebrata; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
 Mus.
 REFERENCE 1 (bases 1 to 337)
 AUTHORS Marra,M., Hillier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,
 Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M.,
 Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,
 Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and
 Waterston,R.
 TITLE The WashU-BHMI Mouse EST Project
 JOURNAL Unpublished (1996)
 COMMENT
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 Washington University School of MedicineP
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
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 Fax: 314 286 1810
 Email: mouseest@watson.wustl.edu
 This clone is available royalty-free through LLNL ; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 MGI:289911
 Seq primer: -28M13 rev2 from Amersham
 High quality sequence stop: 292.
 FEATURES
 source Location/Qualifiers
 1..337
 /organism="Mus musculus"
 /strain="C57BL/6J"
 /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
 polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
 was primed with a Not I - oligo(dT) primer [5'

TGTTACCAATCTGAAGTGGGAGCGGCCGCGAAATTTTTTTTTTTTTTTTTTTTTT
T 3'], on equal amounts of mRNA from 2 13.5dpc and 2
14.5dpc embryos [total RNA provided by Minoru Ko, Wayne
State Univ., from 2]; double-stranded cDNA was ligated to
Eco RI adaptors (Pharmacia), digested with Not I and
cloned into the Not I and Eco RI sites of the modified
pT7T3 vector. Library went through one round of
normalization, and was constructed by Bento Soares and
M.Fatima Bonaldo."

/clone="479167"

/clone_lib="Soares mouse embryo NbME13.5 14.5"

/sex="unknown"

/tissue_type="embryo"

/dev_stage="13.5-14.5dpc total fetus"

/lab_host="DH10B"

<1..>337

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121 aactatggga aaaagagatg aagctcacgg atatccggtt ggaggccctc aactctgccc
181 accagctgga ccagcttcgg gagaccatgc acaatatgca gttggagggtg gacctgctga
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gb|AA049124|AA049124 mj46f04.r1 Soares mouse embryo NbME13.5 14.5 Mus
musculus cDNA clone 479167 5'
Length = 337

Plus Strand HSPs:

Score = 206 (92.9 bits), Expect = 3.9e-19, P = 3.9e-19

Identities = 42/60 (70%), Positives = 51/60 (85%), Frame = +3

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Sbjct: 84 EPEKKEVSELRLSELWEKEMKLTDIRLEALNSAHQLDQLRETMHNMQLVDLLKAENDRLK 263
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LOCUS AA185349 348 bp mRNA EST 07-JAN-1997
DEFINITION mu51c03.r1 Soares mouse lymph node NbMLN Mus musculus cDNA clone
642916 5'.
ACCESSION AA185349
NID g1769059
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
Eukaryotae; mitochondrial eukaryotes; Metazoa; Chordata;
Vertebrata; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Mus.
REFERENCE 1 (bases 1 to 348)

AUTHORS Marra,M., Billier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T., Geisel,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M., Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B., Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and Waterston,R.

TITLE The WashU-BHMI Mouse EST Project

JOURNAL Unpublished (1996)

COMMENT

Contact: Marra M/Mouse EST Project
WashU-BHMI Mouse EST Project
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4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:394908
Seq primer: -28M13 rev2 from Amersham
High quality sequence stop: 336.

FEATURES

source Location/Qualifiers

1..348

/organism="Mus musculus"

/strain="C57BL/6J"

/note="Vector: pT7T3D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; [5'
TGTTACCAATCTGAAGTGGGAGCGCGCGGATCTTTTTTTTTTTTTTTTTTTTTTTT
3']; double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not
I and Eco RI sites of the modified pT7T3 vector. RNA
provided by Dr. Bertrand Jordan. Library constructed and
normalized by Bento Soares and M.Fatima Bonaldo."

/clone="642916"

/clone_lib="Soares mouse lymph node NbMLN"

/sex="male"

/dev_stage="4 weeks"

/lab_host="DH10B"

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BASE COUNT 93 a 95 c 78 g 82 t

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121 ctatctctgc gttcttacca cttgttctgg ccaatagaac tccggatcaa gaggcagaat
181 tcctcagata gcatctccag cctcaacagc atcaccagcc attccagcat cggcagcagc
241 aaagatgctg atgccaagaa gaaaaagaag aagagttggg taagtaaagg cttggagata
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//

gb|AA185349|AA185349 mu51c03.r1 Soares mouse lymph node NbMLN Mus
musculus cDNA clone 642916 5'
Length = 348

Plus Strand HSPs:

Score = 154 (69.4 bits), Expect = 4.4e-12, P = 4.4e-12
Identities = 27/42 (64%), Positives = 40/42 (95%), Frame = +1

Query: 1656 DLRIRRQHSDDSVSSINSATSHSSVGSNIESDSKKKKRKNWL 1697
 +LRI+RQ+SSDS+SS+NS TSHSS+GS+ ++D+KKKK+K+W+
Sbjct: 157 ELRIKRONSSDSISLNSITSHSSIGSSKDADAKKKKKSWV 282

FIGURE 12a

"SIM output with parameters:
substitution scores in BLOSUM62
O = 12, E = 4"

Sequence 1: hu1, 1702 residues
Sequence 2: hu2, 2350 residues

List of local alignments with score >= 100.0

46.8% identity in 1726 residues overlap; Score: 2538.0; Gap frequency: 9.3%

```

hu1,      78 DPESQRRKRTVQNVLDRQNLEETMSSLRGSQVTHSSLEMTCTYDS--DDANPRSVSSLSNR
hu2,     639 DPEARLRRTVKNIADLRQNLEETMSSLRGTQVTHSTLETTFDTNVTTEMSGRSILSLTGR
      *** * *** * ***** * * * * * * * * * * * * * * * * * * * * * * *
hu1,     136 SSPLSWRYGQSSPRLQAGDAPSVGGSCRSEGTPAWYMHGERAHYSHTMPMRSPSKLSHIS
hu2,     699 PTPLSWRLGQSSPRLQAGDAPSMGNGYPPRANASRFINTESGRVYVSAPLRRQLASRGSS
      ***** * * * * * * * * * * * * * * * * * * * * * * *
hu1,     196 RLEL-VESLDSDEVDLKS-----GYMSDSLGMKMTMEDDDITG-----
hu2,     759 VCHVDVSDKAGDEMDLEGISMDAPGYMSDGDVLSKNI-RTDDITSGYMTDGGGLGLYTRRL
      * * * * * * * * * * * * * * * * * * * * * * *
hu1,     235 -----WDESSSISSGLSDASDNLSSSEEFNASSSLNSLP
hu2,     818 NRLPDGMVAVRETQRNTSLGLGDADSWDDSSSVSSGISDTIDNLTDDINTSSSISSYA
      * * * * * * * * * * * * * * * * * * * * * * *
hu1,     268 STPTASRRNSTIVLRDSEKRS LAESGLSWFSESEEKAPKKLEYDSGLKMEPGTSKWRR
hu2,     878 NTPASSRKNLDV--QTDAEKHSQVERNSLWSGDDVKKSDGGS--DSG-IKMEPG-SKWRR
      ** * * * * * * * * * * * * * * * * * * * * * * *
hu1,     328 ERPESCDSSKGGELKKPISLGHPGSLKKGKTPPVAVTSPITHTAQ--SALKVAGK---P
hu2,     932 NPSDVSDSDKSTSGRKNPVISQTGSWRRGMTAQVGITMPRTKPSAPAGALKTPGTGKTD
      * * * * * * * * * * * * * * * * * * * * * * *
hu1,     383 EGKATDKGKLAVKNTGLQRSSSDAGRDRLSDAKKPPSGIARPSTSG--SFGYKKPP-PAT
hu2,     992 DAKVSEKGR LSPKASQVRSPSDAGRSSGDESKKPLPSSRTPTANANSFGFKKQSGSAA
      * * * * * * * * * * * * * * * * * * * * * * *
hu1,     440 GTATVMQTG-----GSATLSKIQKSSGIPVKPVNGRKTSLDVSN SAE PGFLAPGARSNIQ
hu2,    1052 GLAMITASGVTVTSRSATLGKIPKSSAL-VSR SAGRKSSMDGAQNQDDGYLALSSRTNLQ
      * * * * * * * * * * * * * * * * * * * * * * *
hu1,     495 YRSLPRPAKSSSMSVTGGRGGPRPVSSSIDPSLLSTKQGGLTPSRLKEPTKVASGRTPA
hu2,    1111 YRSLPRPSKNSNRNGAGNRSS-----TSSID-SNISSKSAGLPVKLREPSKTALGSSSLPG
      ***** * * * * * * * * * * * * * * * * * * * * * * *
hu1,     555 PVNQTDREKEKAKAKAVALDSNISLSKISGPESTPKNQASHPTATKLAELPPTPLRATA
hu2,    1166 LVNQTDKEKGISSDNESVASCNSVKVNPAAQPVSSPAQTS LQPGAKYPDVASPTLRLFLG
      ***** * * * * * * * * * * * * * * * * * * * * * * *

```


hu1, 615 RSVKPPSLANLDKV-NSNSLDLPSSSDTTHAS--KVPDLBATSSASGGPL-----P
hu2, 1226 GKPTKQVPIATAENMKNSVVISNPBATMTQQGNLDSPPSGSVLSSGSSSPLYSKNVDLNQ
* * * * *

hu1, 664 SCFTSPAPILNINSASFQGLEMSGFSVPKETRMYPKLSGLHRSMESLQMPMS---LP
hu2, 1286 SPLASSPSSAHSAPSNLTWTGNASSSSAVSKDGLGFQSVSSSLHTSCESIDISLSSGGVP
* * * * *

hu1, 721 SAFPSSTPVPTPPAPPAAP-TEETEELTWSGSPRAGQLDSNQ-----RD
hu2, 1346 SHNSSTGLIASSKDDSLTPFVRTNSVKTTLSSESPLSSPAASPKFCRSTLPRKQSDPHLD
* * * * *

hu1, 765 RNTLPKKGLRY----QLQSQEETKRERHSHTIGGLPESDDQSELSPPALPMSLSAKGQL
hu2, 1406 RNTLPKKGLRYTPTSQRLTQEDAKEWLRSHSAGGLQDTAANSFPSSGSSSVTSPSGTRFNF
***** * * * *

hu1, 821 TNIVSPTAAT-----TPRITRSNSIPTHEAAFELYSGSQM-GSTLSLAERPCKMIRSGSF
hu2, 1466 SQLASPTTVTQMSLSNPTMLRTHSLSNADGQYDPTDSRFRNSSMSLDEKSRTMSRSGSF
* * * * *

hu1, 875 RDPTDDVHGSVLSLASSASSTYSAAERMQSEQIRKLRELESSQEKVATLTSQLSANAN
hu2, 1526 RDGFEEVHGSSLSLVSSSTLSVYSTPEEKQSE-IRKLRELDASQEKVSALTQTANAE
* * * * *

hu1, 935 LVAAFEQSLVNMTSRLRHLAETAEEKDTELLDLRETIDFLKKKNSEAAVIGQALNASET
hu2, 1585 LVAAFEQSLGNMTIRLQSLTMTAEQKDSLELNRKTIELLKKQNAQAINGVINTPEL
***** * * * *

hu1, 995 TPK-----ELRIKRONSSDISSLNSITSHSSIGSSKDADAKKKKKSWVYELRSSF
hu2, 1645 NCKGNGTAQSADLRIRRHQSSDSVSSINSATSHSSVGSNIESDSKKKKRKNW---LRSSF
* * * * *

hu1, 1047 NKAFSIRKGPKSASSYSIDIEIATPDSSAPSSPKLQHGSTETASPSIKSSTLSSVGTDTV
hu2, 1702 KQAFGKKKSPKSASSHSDIE--TTDSSLPSSPKLPHNGSTGTPLLRNHSNSL-----
* * * * *

hu1, 1107 EGPABPAPHTRLFHANEETPEKKEVSELRLSELWEKEMKLTDIRLEALNSAHQLDQLRET
hu2, 1755 -----ISECMDSEAEVTVMQLRNELRDKEMKLTDIRLEALNSAHQLDQLREA
* * * * *

hu1, 1167 MBNMQLEVDDLKAENDRLKVAPGPSSGSTPGQVPGSSALS-SPRRSLGLALTHSFGPSLA
hu2, 1801 MNRMQSEIEKLKAENDRLK---SESQSGGCSRAPSQVSIASPRQSMGLS-QHSLNLTES
* * * * *

hu1, 1226 DTDLSPMDGISTCGPKEEVT--LRVVVRMPQHI IKGDLKQOEFFLGCSKVSGKVDWKML
hu2, 1857 TSLDMLDDTGECSARKEGGRHVKIVVSFQEEMKWKEDSRPHLFLIGCIGVSGKTKWDVL
* * * * *

hu1, 1284 DEAVFQVKDYISKMDPASTLGLSTESIBGYSISHVKRVLDAEPPMPPCRGRVNN---I
hu2, 1917 DGVVRRLFKEYIIHVDPVSQLGLNSDVLGYSIGEIKRSNTSETPELLPCGYLVGENTTI
* * * * *

```
hu1,      1341 SVSLKGLKREKCVDSLVEFETLIPKPMQBYISLLKHRRVLVSGPSGTGKTYLTNRLEAYL
hu2,      1977 SVTVKGLAENSLDSLVEFSLIPKPIQRYVSLIEHRRILSGPSGTGKTYLANRLSEYI
          ** *** * ***** * * * * * * * * * * * * * * * * * * * *
hu1,      1401 VERSGREVTEGIVSTFNMBQQSCDKLQLYLSNLANQIDRETGIGDVPLVILLDDLSEAGS
hu2,      2037 VLREGRELTGVIATFNVDBKSSKELRQYLSNLADQCENSENNAVDMPLVIILDNLHHVSS
          * * * * * * * * * * * * * * * * * * * * * * * * * *
hu1,      1461 ISELVNGALTCKYHKCPYIIGTTNQPVKMTPNHGLHLSFRMLTFSNNVEPANGFLVRYLR
hu2,      2097 LGEIFNGLLNCKYHKCPYIIGTMNQATSSTPNLQLHHNFRWVLCANHTEPVKGFLGRFLR
          * * * * * * * * * * * * * * * * * * * * * * * * * *
hu1,      1521 RKLVESDS DINANKEELLRVLDWVFKLWYHLTFLEKHSSTDFLIGPCFFLSCPIGIEDF
hu2,      2157 RKLME TEISGRVRNME LVKIIDWIPKVWHHLNRFLEAHSSSDVTIGPRLFLSCPIDVDGS
          *** * * * * * * * * * * * * * * * * * * * * * * *
hu1,      1581 RTWFIDLWNSIIPYLOEGAKDGKIVHGQKAAWEDPVEWVRDTLPWPSAQDQS--KLYH
hu2,      2217 RVWFTDLWNYSIIPYLL EAVREGQLYGRAPWEDPAKWVMDTYPWAAS PQHEWPPLLQ
          * * * * * * * * * * * * * * * * * * * * * * * * *
hu1,      1639 LPPPTVGPBSIASPPEDRTVKDSTPSSLDSDPLMAMLLKLQEAANY
hu2,      2277 LRPEDVGFDGYSMPREGSTSKQMPPSDAEGDPLMNMLMRLQEAANY
          * * * * * * * * * * * * * * * * * * * * * * * *
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WARNING: 49 local alignments have not been reported because of score < 100.0

"SIM output with parameters:
substitution scores in BLOSUM62
O = 12, E = 4"

Sequence 1: Cel, 1583 residues
Sequence 2: hu2, 2350 residues

List of local alignments with score >= 54.0

32.8% identity in 504 residues overlap; Score: 490.0; Gap frequency: 6.9%

```
Cel,      1058 VIELKQELKERDSALYEVRLDNLDRAREVDVLRET VNKLTENKQLKKEVDKLTNGPATR
hu2,      1766 VMQLRNELRDKEMKLTDIRLEALSSAHQLDQLREAMNRMQSEIEK LKAENDRLKSESQGS
          * * * * * * * * * * * * * * * * * * * * * * * *
Cel,      1118 ASSRASIPVIYD-----DEHVDAA CSST-----SASQSSKRSSGCNSIKVTNVV
hu2,      1826 GCSRAPSQVSISASPRQSMGLSQHSLNLTESTSLDMLDDTGECSARKEGGRHV KIVVSF
          *** * * * * * * * * * * * * * * * * * *
Cel,      1163 DIAGEISSIVNPDKEIIVGYLAMSTSQSCWKDIDVSI LGLFEVYLSRIDVEHQLGIDARD
```

hu2, 1886 QEEMKWKEDSRPHL-FLICIGVS-GKTKWDVLDGVVRRLFKEYIIHVDPVSQGLGNS-D
* * * * *
Cel, 1223 SILGYQIGELRRVIGDSTMTITSHPTDILTSSSTIRMFHMGAAQSRVDSLVDMLLPKQM
hu2, 1943 SVLGYSIGEIKRSNTSETPELLPCGY-LVGENTTISVTVKGLAENSLDSLVPESLIPKPI
* * * * *
Cel, 1283 ILQLVKSILTERRLVLGATGIGKSKLAKTLAAYVSIRTNQ--SEDSIVNISIPENKKEE
hu2, 2002 LQRYVSLLEHRRRIILSGPSGTGKTYLANRLSEYIVLREGRELTGVIATFNVDKSSKE
* * * * *
Cel, 1341 LLOVERRLEKIILRSKESCI-----VILDNIPKNRIAFVVSFANV-PLQNEGPFVVCVCTV
hu2, 2062 LRQYLSNLADQCSENNAVDMPLVIILDNL--HBVSSLGEIFNGLLNCKYHKCPYIIGTM
* * * * *
Cel, 1395 NRY--QIPELQIHHNFKMSVMSNRLE---GFILRYLRRRAVEDEYRLTVQMPSELFRKIID
hu2, 2120 NQATSSTPNLQLHHNFRWVLCANHTEPVKGFLGRFLRKLMEITEISGRVRN-MELVKIID
* * * * *
Cel, 1450 FFPIALQAVNNFIEKTSVDVTVGPRACLNCPLTVDGSREWFIRLWNNFIPYLERVARD
hu2, 2179 WIPKVVHHLNRFLEAHSSSDVTIGPRLFLSCPIDVDGSRVWFDTLWNYSIIPYLLEAVRE
* * * * *
Cel, 1510 GKKTFGRCTSFEDPTDIVSEKWPW
hu2, 2239 GLQLYGRRAFWEDPAKWVMDTYPW
* * * * *

35.5% identity in 112 residues overlap; Score: 165.0; Gap frequency: 1.8%

Cel, 11 IYTDWANRHLKSGSLKXSIRDISNDFRDYRLVSQLINVIVPINEFSPAFTKRLAKITSNL
hu2, 11 IYTDWANHYLTKSGHKRLIKDLQODVTDGVLLAQIIQVVA--NEKIEDINGCPKNRSQMI

Cel, 71 DGLETCLDYLKNLGLDCSKLTKTDIDSGNLGAVLQLLFLLSTYKQKLRQLKK
hu2, 69 ENIDACLNFLAAKGINIQGLSAEEIRNGNLKAILGLFFSLSRKQOQQQPQK
* * * * *

24.8% identity in 163 residues overlap; Score: 80.0; Gap frequency: 3.7%

Cel, 877 GSQSLSLASTT--AYGSLNEKYEHAIKRDMDARDLECYKNTVDLSLTKKQENYGALFDLFEQKL
hu2, 1534 GSSLSLVSTLSVYSTPEEKQSEIRKLRLRELDASQEKVSALTTQLTANAHLVAAFEQSL
* * * * *
Cel, 935 RKLTOHIDRSNLKPEEAIRFQDIAHLRDISNHLASNSAHANEGAGELLRQPSLESVASH
hu2, 1594 GNMTIRLQSLTMTAEQK---DSELNELRKTIELLLKKQNAQAQAAINGVINTPELNCKGNG
* * * * *
Cel, 995 RSSMSSSSKSSKQEKISLSSFGK-NKKSWIRSSLSKFTKKKNK
hu2, 1651 TAQSADLRIRRHSSDSVSSINSATSHSSVGSNIESDSKKKKR
* * * * *

58.6% identity in 31 residues overlap; Score: 74.0; Gap frequency: 6.5%

F
Cel, 653 GYPDNFEDSSSLSSGISDNNELDDISTDDL
hu2, 840 GDADSWDDSSSVSSGISDT--IDNLSTDDIN
* * * * * * * * * *

42.9% identity in 60 residues overlap; Score: 64.0; Gap frequency: 6.7%

Cel, 984 RQPSLESVASHRSMSSSSKSKQEKISLSSFGKNKSWIRSSLK-FTRKKKNKYDEAH
hu2, 1661 RQHSSDSVSSINSATSHSSVGS---NIESDSKKKKRKNWLRSSFKQAFGKKKSPKSASSH
* * * * * * * * * * * * * * * * * * * *

22.0% identity in 91 residues overlap; Score: 56.0; Gap frequency: 0.0%

Cel, 140 SKLPSPRVATSATASATNPNSNFPQMSTSRLOTQSRISKIDSSKIGIKPKTSGLKPPSS
hu2, 177 SRLSGPTARVSAAGSEAKTRGGSTANNRRSQSFNNYDKSKPVTSPPPPPSSHEKEPLAS
* * * * * * * * * * * * * * * * * * *

Cel, 200 STTSSNNTNSFRPSSRSSGNNNVGSTISTSA
hu2, 237 SASSHPGMSDNAPASLESGSSSTPTNCSTSS
* * * * * * * * * *

WARNING: 44 local alignments have not been reported because of score < 54.0

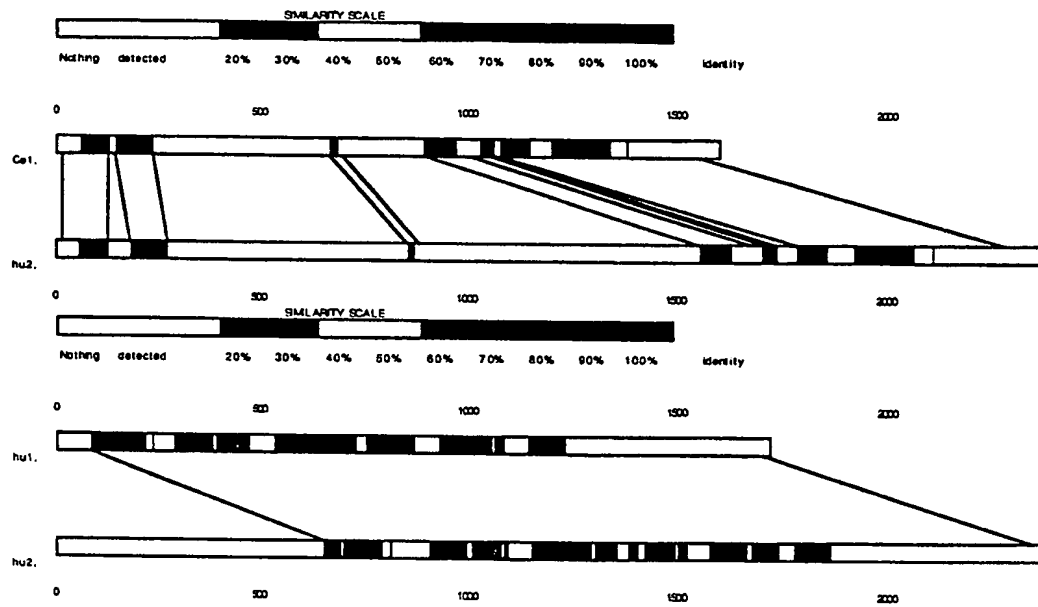


Figure 12b

Tuesday, 18 November 1997 10:09

fig 13 pCB201 (1 > 5082) Site and Sequence

Enzymes : 100 of 148 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

Page

fig 13 pCB201

GACGGATCGGGAGATCTCCCGATCCCTATGGTCGACTCTCAGTACAATCTGCTCTGATGCCGCATAGTTAAGCCAGTATCTGCTCCCTGCTTGTGTGT
CTGCTAGCCCTCTAGAGGGCTAGGGGATACCACTGAGAGTCATGTTAGACGAGACTACGGCGTATCAATTGGTTCATAGACGAGGGACGAACACACAA
T D R E I S R S P M V D S Q Y N L L . C R I V K P V S A P C L C V

GGAGGTCGCTGAGTAGTCGCGAGCAAAATTTAAGCTACAACAAGGCAAGGCTTGACCGACAATTGCATGAAGAATCTGCTTAGGGTTAGGCGTTTGGG
CCTCCAGGACTCATCAGCGCTCGTTTTAAATTCGATGTTGTTCCGTTCCGAATGGCTGTTAAGCTACTTCTTAGACGAATCCCAATCCGCAAAACGG
G G R . V V R E O N L S Y N K A R L D R O L H E E S A . G . A F C

CTGCTTCGCGATGACGGGCGAGATATACGCGTTGACATTGATTATTGACTAGTTATTAATAGTAATCAATTACGGGGTCATTAGTTCATAGCCCATATA
GACGAAGCGCTACATGCCGGTCTATATGCGCAACTGTAACATAAATGATCAATAATTATCATTAGTTAATGCCCGAGTAATCAAGTATCGGGTATAT
A A S R C T G O I Y A L T L I I D . L L I V I N Y G V I S S . P I Y

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ACCTCAAGGCGCAATGATTGAATGCCATTTACCGGGCGGACCGACTGGCGGGTGTCTGGGGGGGGTAACGTCAGTTATTACTGCATACAAGGGTATCA
G V P R Y I T Y G K V P A W L T A Q R P P P I D V N N D V C S H S

AACGCCAATAGGGACTTTCCATTGACGTCAATGGGTGGACTATTTACGGTAACTGCCACTTGGCAGTACATCAAGTGATCATATGCCAAGTACGCC
TTGCGGTATTCCTGAAAGGTAAGTCACTGACGTACCCACCTGATAAATGCCATTTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTTATGCGGG
N A N R D F P L T S M G G L F T V N C P L G S T S S V S Y A K Y A

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GGATAACTGCAGTTACTGCCATTTACCGGGCGGACCGTAATACGGGTGATGACTGGAATACCTGAAAGGATGAACCGTCATGTAGATGCATAATCAG
P Y . R O . R . M A R L A L C P V H D L M G L S Y L A V H L R I S H

TCGCTATTACCATGGTGATGGGTTTTGGCAGTACATCAATGGCGGTGGATAGCGGTTTGACTCAGGGGATTTCAGGTCTCCACCCCATGACGTCAA
AGCGATAATGGTACCCTACGCCAAAACCGTCACTGAGTTACCGGCACCTATCGCCAAACTGAGTGCCCTAAAGGTTACAGAGGTGGGGTAAGTGCAGT
P Y Y H G D A V L A V H O W A V I A V . L T G I S K S P P H . R Q

TGGGAGTTTGTGTTGGCACCAAAATCAACGGGACTTTCCAAAATGTCGTAACAAC TCCGCCCATGACGCAATGGGCGGTAGCGGTGACGTTGGGAG
ACCTCAAAACAAAACCGTGTGTTTGTGCTGAAAGGTTTACAGCATGTTGAGGCGGGGTAAC TGCTTTACCGGTCATCCGACATGCCACCTC
W E F V L A P K S T G L S K M S . Q L R P I D A N G R . A C T V G

GTCATATTAAGCAGAGCTCTCTGGCTAACTAGAGAACCCTACTGCTTACTGGCTTATCGAAATTAATACGACTCACTATAGGGAGACCAAGCTGGCTAGC
CAGATATATTGCTCGAGAGACCGATTGATCTCTGGGTGACGAATGACCGAATAGCTTTAATTATGCTGAGTGATATCCCTCTGGGTTCGACCGATCG
G L Y K G S S L A N . R T H C L L A Y R N . Y D S L . G D P S V L A

GTTTAACTTAAGCTTACCATGGGGGTTCTCATCATCATCATCATGGTATGGCTAGCATGACTGGTGGACAGCAATGGGTCGGGATCTGTACGAG
CAATTTGAATTCGAATGGTACCCCCAAGAGTAGTAGTAGTAGTACCATACCGATCGTACTGACCACTGTGCTTTACCCAGCCCTAGACATGCTG
F K L K L T M G G S H H H H H G M A S M T G G Q Q M G R D L Y D

T7 promoter priming site

ProBond binding domain

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GATGACGATAAGGTACCTAGGATCCATATGCCCTCTTGGCGTCGAGGTGTCAATAACATATCAGTCTCCCTCAAAGGCTGAAGGAGAAATGCGTCGACAT
CTACTGCTATTCCATGGATCCTAGGTATACGGAGGAACGGCAGCTCCACATTTATGTATAGTCAGAGGGAGTTTCCAGACTTCTCTTTACGCAGCTG
pCB201 insert = U4
U4 ORF
D D D K V P R I H M P P C R R G V N N I S V S L K G L K E K C V D
GCC TGGTGTTCGAGACGCTGATCCCCAAGCCGATGATGCAGCACTACATAAASCTCTGCTGAAGCACCGGCGCCTCGTCTCTCGGGCCCCAGCGGAC
CGGACCACAAGCTCTGCGACTAGGGGTTCTGGCTACTACGTCGTGATGTATTCGGAGGACGACTTCGTGGCCGCGGAGCAGGAGAGCCCCGGGTCGCCGTS
pCB201 insert = U4
U4 ORF
S L V F E T L I P K P M M Q H Y I S L L L K H R R L V L S G P S G T
GGGCAAGACCTACCTGACCAATCGCTTGGCCGAGTACCTGGTGGAGCGCTCTGGCCGTGAGGTACAGAGGGCATCGTCAGCACCTTCAACATGCACCA
CCCCTTCGGATGGACTGGTTAGCGAACC GGCTCATGGACCACCTCGCGAGACCGGCACTCCAGTGTCTCCCGTAGCAGTCGTGGAAGTTGTACGTGGT
pCB201 insert = U4
U4 ORF
G K T Y L T N R L A E Y L V E R S G R E V T E G I V S T F N M H Q
CASTCTTGCAAGGATCTGCAACTGTATCTTTCCAACCTAGCCAACCAAGATACACCGGGAACAGGAATTGGGGATGTGCCCTGGTGATTCTATTGGATG
GTCAGAACGTTCTAGACGTTGACATAGAAAGGTTGGATCGGTTGGTCTATCTGGCCCTTTGTCTTAACCCCTACACGGGGACCAC TAAGATAACCTAC
pCB201 insert = U4
U4 ORF
D S C K D L Q L Y L S N L A N Q I D R E T G I S D V P L V I L L D
ACCTGAGTGAAGCAGGCTCCATCAGTGAGTTGGTCAATGGGGCCCTCACCTGCAAGTATCATAAATGTCCC TATATTATAGGTACCAACCAATCAGCTG
TGGACTCACTTCGTCCGAGGTAGTCACTCAACCAAGTTACCCCGGGAGTGGACGTTTATAGTATTTACAGGATATAATATCCATGGTGGT TAGTGGAC
pCB201 insert = U4
U4 ORF
D S E A G S I S E L V A G A L T C K Y H K C P Y I I G T T N Q P V
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TTTTTACTGTGGGTTGGTACCGAACGTGAACCTGSAAGTCTACAAC TGAAGAGGTTGTTGCACCTCGGTGGTTACCGAAGGACCAAGCAATGGACTCC
pCB201 insert = U4
U4 ORF
K N T P N H G L H L S F R M L T F S N N V E P A N G F L V R Y L F

BASE COUNT 173 a 168 c 141 g 128 t
ORIGIN

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421 gctagtcaat gcccgactg ctttacgcc cccgcagcct ccagttcag gacctagtga
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DEFINITION fa05f06.s1 Zebrafish ICRFzfls Danio rerio cDNA clone 5D16 3'.
ACCESSION AA495042
NID g2225470
KEYWORDS EST.
SOURCE zebrafish.
ORGANISM Danio rerio
Eukaryotae; mitochondrial eukaryotes; Metazoa; Chordata;
Vertebrata; Actinopterygii; Neopterygii; Teleostei; Euteleostei;
Ostariophysi; Cypriniformes; Cyprinoidea; Cyprinidae; Rasborinae;
Danio.
REFERENCE 1 (bases 1 to 418)
AUTHORS Clark,M., Lehrach,H., Johnson,S., Marra,M., Eddy,S., Billier,L.,
Allen,M., Bowles,L., Dubuque,T., Geisel,G., Jost,S., Kucaba,T.,
Lacy,M., Le,N., Lennon,G., Martin,J., Moore,B., Schellenberg,K.,
Steptoe,M., Tan,F., Theising,B., White,Y., Wylie,T., Waterston,R.
and Wilson,R.
TITLE WashU Zebrafish EST Project
JOURNAL Unpublished (1997)
COMMENT
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Tel: 314 286 1800
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Steve Johnson lab internal ID - P2_60 NOTE - For this library, the
CLONE id field represents a position identifier on the original
cDNA library preparation plate. cDNA Library Preparation: Matthew
Clark. cDNA Library Arrayed by: Matthew Clark. DNA Sequencing by:
Washington University Genome Sequencing Center Clone distribution:
Genome Systems, St. Louis, and Max Planck Institut fuer Molekulare
Genetik, Berlin Tel +49 30 84 13 1235
Seq primer: -40ml3 ET from Amersham
High quality sequence stop: 416.
FEATURES
source Location/Qualifiers
1..418
/organism="Danio rerio"
/note="Vector: pSPORT1; Site_1: NotI; Site_2: SalI; 1st
strand cDNA was primed with a Not I - oligo(dT)15 primer
[5'pGACTAGTTCTAGATCGCGAGCGCCGCCCTTTTTTTTTTTT3'], on
mRNA from pooled 26 somite zebrafish embryos;
double-stranded cDNA was ligated to Sal I adaptors (BRL),
digested with Not I and cloned into the Not I and Sal I
sites of the pSPORT1 vector (BRL). Library was constructed
by Matthew Clark (Lehrach lab; ICRF, London and Max
Planck Institut fuer Molekulare Genetik, Berlin) and was
not biochemically normalised. 70,000 clones from this
library were arrayed on high density filters and
subsequently screened by oligonucleotide hybridization
fingerprinting to identify unique or minimally redundant
clones for more intensive analysis."

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AGGAAGCTGGTAGAGTCAGACAGCGACATCAATGCCAACAGGAAGAGCTGCTTCGGGTGCTCGACTGGGTACCCAAGCTGTGGTATCATCTCCACACCT
TCCTTCGACCATCTCAGTCTGTGCTGTAGTTACGGTTGTTCTTCGACGAAGCCACGAGCTGACCCATGGGTTCGACACCATAGTAGAGGTGTGGG

pCB201 insert = U4

U4 ORF

R K L V E S D S D I N A N K E E L L R V L D V V P K L V Y H L H T

TCCTTGAGAAGCACAGCACCTCAGACTTCCTCATCGGCCCTTGCTTCTTCTGTGCTGTCCCATTGGCATTGAGGACTTCGGGACCTGCTTCATTGACCT
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pCB201 insert = U4

U4 ORF

F L E K H S T S D F L I G P C F F L S C P I G I E D F R T V F I D L

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CACCTTGTGAGATAGTAAGGGATAGATGTCCTTCCTCGGTTCCTACCTATTTCAGGTACCTGTCTTCGACGAACCTCCTGGGTCACCTTACCCAG

pCB201 insert = U4

U4 ORF

V N N S I I P Y L Q E G A K D G I K V H G Q K A A W E D P V E W V

CGGGACACACTTCCTTGGCCATCAGCCCAACAAGACCAATCAAGCTGTACCACCTGCCCCACCCACCGTGGGCCCTCACAGCATTGCTCAGCTCCG
GCCCTGTGTGAAGGGACCGGTAGTCGGGTTGTTCTGGTTAGTTTCGACATGGTGGACGGGGTGGGTGGCACCCGGGAGTGTCTGAACGGAGTGGAGGGG

pCB201 insert = U4

U4 ORF

R D T L P W P S A Q Q D Q S K L Y H L P P P T V G P H S I A S P P

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pCB201 insert = U4

U4 ORF

E D R T V K D S T P S S L D S D P L M A M L L K L Q E A A H Y I E S

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AGSTCTAGCTCTTTGGTAGGACCTGGGGTTGGAAGTCGGTTGTGAATTCCTAAGCCGTTAGTGACAGTGGGGGCCGTGCTCTTGGACCGTAGTGGG

pCB201 insert = U4

U4 ORF

P D F E T I L D P N L O A T L G F G N H C H P R T A E R W H Q L

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Page

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AGAATCGAGGAGGAGAGGGGAGAGGAGAAAGTCTCGTGACCGAGAGGTGCGGGTCTCTCTTGCTCCTCCTCTCTACTTTCTCCTCCTGTCCAA 2300

pCB201 insert = U4

S . L L L S P L L F Q S T G S P A P G G E Q E G G G D E R G G T G

CTTGGTGCTGTACCTTTGAGAACTTCCTAGGAAGGAATGGTGGGGTGGCGTTTGGGAACTTGTCCTCTTAAACACATTTACTGGCCTCCTCTAATGACT
GAACCACGACATGGAACTCTTGAAGGATCTCTCTTACCACCCACCGCAACCCCTTGAACACGGGGGATTTGTGTAATGACCGGAGGAGATTACTGA 2400

pCB201 insert = U4

S V C C T F E N F L G R N G G V A F G N L C P L N T F T G L L . . L

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AACCCTTTTCTACTAAGACCCAGAAAGGAACTGAAGAACAAGTTAATGTTTGGAGACCGAAAGACCCCTCCCAAGTCTTTGTAGTTTGTGAGC 2500

pCB201 insert = U4

V G K D D S G S F P . L L V S I T N S V A F V G G V Q K T S K H C

AGCAGTTCTTAATGATTCTACAAGCAACCTGAGAGAGACAGTCTTGTGAGGGAGATCTGGGGAGGCGAGGAAGCTCCTCAGATTTTCTCACAGACCC
TCGTCAAGGATTTACTAAGAGTGTTCTGTTGGGACTCTCTCTGTGACAACTCCCTCTAGACCCCTCCGTCCTTCGAGGAGTCTAAAAGAGTGCTGGG 2600

pCB201 insert = U4

S S S . M I L T S N P E R D S L V R E I V G R Q E A P O I F S Q T

TTCCCAATTCATCACCACGCAACAC TCGTCCGGAATTCGTGAGATATCCAGCACAGTGGCGGCCGCTCGAGTCTAGAGGGCCGTTTAAACCCGCTG
AAGGTTAAGGTAGTGGTGACGGTTGTGAGCAGCCTTAAGACGCTATAGSTCGTGTACCGCCGGCGAGCTCAGATCTCCCGGGCAAAATTGGGGCAG 2700

pCB201 insert = U4

L P N S I T T A N T R P E F C R Y P A Q V R P L E S R G P V . T R .

ATCAGCTCGACTGTGCCTTCTAGTTGCCAGCCATCTGTTGTTTGCCTTCCCCGTCCTTCTTGACCTTGGAGGTGCCACTCCCACTGTCTTTCC
TAGTCGGAGCTGACACGGAAGATCAACGGTCGGTAGACAACAACGGGGAGGGGACGGAAGGAAC TGGGACCTTCCACGGTGAGGGTGACAGGAAAGG 2800

S A S T V P S S C Q P S V V C P S P V P S L T L E G A T P T V L S

TAATAAATGAGGAAATGCATCGCATTGTCTGAGTAGGTGTCATTCTATTCTGGGGGGTGGGGTGGGGCAGGACAGCAAGGGGGAGGATTGGGAAGACA
ATTATTTTACTCTTTAACGTAGCGTAACAGACTCATCCACAGTAAGATAAGACCCCCACCCACCCCGTCTGTCTGTTCCCCCTCCTAACCTTCTGT 2900

. . N E E I A S H C L S R C H S I L G G G V G Q D S K G E D V E D

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TATCGTCCGTACGACCCCTACGCCACCCGAGATACCGAAGACTCCGCTTTCTTGGTCGACCCGAGATCCCCCATAGGGGTGCGCGGGACATCGCGGGS 3000

N S R H A G D A V G S M A S E A E R T S V G S R G Y P H A P C S G A

ATTAAGCGCGGGGGTGTGGTGGTTACGCGCAGCGTGACCGCTACACTTGCAGCGCCCTAGCGCCGCTCTTTCGCTTTCTTCCCTTCTTCTCGG
TAATTCGCGCGCCACACCAATGCGGTGCGCTGCGGATGTGAACGCTCGGGGATCGCGGCGAGGAAAGCGAAAGGAAGGAAAGAGAGCGS 3100

L S A A G V V V T R S V T A T L A S A L A P A P F A F F P S F L A

ACGTTCCCGGCTTTCCCGTCAAGCTCTAAATCGGGCATCCCTTTAGGGTTCGATTTAGTGCTTACGGCACCTCGACCCCAAAAACCTTGATAGG
TGCAAGCGCGGAAAGGGCAGTTCGAGATTTAGCCCGTAGGGAATCCCAAGGCTAAATCAGAAATGCCGTGGAGCTGGGGT TTTTGAAC TAATCC 3200

T F A G F P R Q A L N R G I P L G F R F S A L R H L D P K K L D .

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CCTACCAAGTGCATCACCCGGTAGCGGACTATCTGCCAAAAAGCGGAACTGCAACCTCAGGTGCAAGAAATTATCACCTGAGAACAAAGTTTGACG
G D G S R S G P S P . . T V F R P L T L E S T F F N S G L L F Q T G
AACAACACTCAACCTATCTCGGTCTATTCTTTGATTATAAGGGATTTGGGGATTTTCGGCTATTGGTTAAAAATGAGCTGATTTAACAAAAATTT
TTGTTGTGAGTTGGGATAGAGCCAGATAAGAAAC TAAATATTCCTTAAACCCCTAAAGCCGGATAACCAATTTTTTACTCGACTAAATTTGTTTTAA
T T L N P I S V Y S F D L . G I L G I S A Y V L K N E L I . Q K F
AACGCGAATTAATTCGTGGAATGTGTGTCAGTTAGGGTGTGGAAAGTCCCAGGCTCCCCAGGCAGGCAGAAGTATGCAAAGCATGCATCTCAATTAGT
TTTCGCTTAATTAAGACACCTTACACACAGTCAATCCACACCTTTTCAGGGGTCCGAGGGGTCCGTCCTTCATACGTTTCGTACGTAGAGTTAATCA
N A N . F C G M C V S . G V E S P Q A P Q A G R S M Q S M H L N .
CAGCAACCAGGTGTGGAAAGTCCCAGGCTCCCCAGGCAGGCAGAAGTATGCAAAGCATGCATCTCAATTAGTACGCAACCATAGTCCCGCCCTAACTCC
GTCGTTGGTCCACACCTTTTCAGGGGTCCGAGGGGTTCGTCCTTCATACGTTTCGTACGTAGAGTTAATCAGTCGTTGGTATCAGGGCGGGGATTGAGG
S A T R C G K S P G S P A G R S M Q S M H L N . S A T I V P P L T P
GCCATCCCCCCCTAACTCCGCCAGTTCCGCCCATTCGCCCCATGGCTGACTAATTTTTTTTATTTATGCAGAGGCCGAGGCCGCTCTGCCTCT
CGGGTAGGGCGGGGATTGAGGCGGGTCAAGGCGGGTAAAGGCGGGGTACCGACTGATTAAAAAAATAAATACGTCCTCCGGCTCCGGCGGAGACGGAGA
P I P P L T P P S S A H S P P H G . L I F F I Y A E A E A A S A S
GAGCTATTCAGAAGTAGTGAAGGAGGCTTTTTTGAGGCGCTAGGCTTTTGCAAAAGCTCCCGGGAGCTTGATATCCATTTTCGGATCTGATCAAGAGA
CTCGATAAGGTCTTCATCACTCTCCGAAAAAACCCTCGGATCCGAAACGTTTTTCGAGGGCCCTCGAACATATAGGTAAAGCCCTAGACTAGTTCTCT
E L F Q K . . G G F F G G L G F C K K L P G A C I S I F G S D Q E
CAGGATGAGGATGTTTCGATGATTGAACAAGATGGATTGCACGCAGGTTCTCCGGCCGCTTGGGTGGAGAGGCTATTCGGCTATGACTGGGCACAACA
GTCTACTCTAGCAAGCGTACTAAGTTGTTCTACCTAACGTGCGTCCAAGAGGCGGGCAACCCACCTCTCCGATAAGCCGATCTGACCCGTGTTG
T G . G S F R M I E Q D G L H A G S P A A V V E R L F G Y D W A Q Q
GACAATCGGCTGCTCTGATGCCGCCGTGTTCCGGCTGTGAGCGCAGGGGCGCCGGTCTTTTTTGTCAGACCGACCTGTCCGGTGCCCTGAATGAAGT
CTTTAGCCGACGAGACTACGGCGGCACAAGGCCGACAGTCGCTCCCGCGGGCCAAGAAAAACAGTTCTGGCTGGACAGGCCACGGGACTTACTTGAC
T I G C S D A A V F R L S A Q G R P V L F V K T D L S G A L N E L
CAGGACGAGGACGCGGGCTATCGTGGCTGGCCACGACGGGCTTCCTTGCGCAGCTGTGCTCGACGTTGTCACTGAAGCGGGAAAGGACTGGCTGCTAT
GTCTGCTCCGTCGCGCCGATAGCACCGACCGGTGCTGCCCGCAAGGAACGCTGACACGAGCTGCAACAGTGAATTCGCCCTTCCTGACCGACGAT
G D E A A R L S V L A T T G V P C A A V L D V V T E A G R D V L L
TGGGCGAAGTGCCGGGCGAGGATCTCTGTGATCTACCTTTGCTCTTGCGGAGAAATATCCATCATGGCTGATGCAATGCGGCGGGTGCATACGCTTGA
ACCGGCTTCACGGCCCCGTCTAGAGGACAGTAGAGTGAACGAGGACGGCTCTTTCAAGGTAGTACCGACTACGTTACGGCGCGGACGATGCGGAC
L S E V P G Q D L L S S H L A P A E K V S I M A D A M R R L H T L D
TCGGGCTACCTGCCCATTCGACCACCAAGCGAAACATCGCATCGAGCGAGCAGTACTCGGATGGAAGCCGGTCTTGTCGATCAGGATGATCTGGACGAA
AGGCGGATGGACGGTAAAGCTGGTGGTTGCTTTGTAGCGTAGCTCGCTCGTGCATGAGCTTACCTTCGGCCAGAACAGCTAGTCTCTACTAGACCTGCTT
P A T C P F D H O A K H R I E R A R T R M E A G L V D Q D D L D E
GAGCATCAGGGGCTCGCGCCAGCCGAACGTTCGCCAGGCTCAAGGCGGCGATGCCCGACGGCGAGGATCTCGTCTGAGCCCATGGCGATGCTGCTTGG
CTGCTAGTCCCCGAGCGCGGTGGCTTGACAAGCGGTCCGAGTTCCGCGGTCAGGCTGCCGCTCTAGAGCAGCACTGGGTACCGCTACCGACGAAGC
E H G G L A P A E L F A R L K A R M P D G E D L V V T H G D A C L

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fig 13 pCB201 (1 > 5082) Site and Sequence

Page

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P N I M V E N G R F S G F I D C G R L G V A D R Y Q D I A L A T R D
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K G W A S E S F S G T P A G . S S S A G I S C V S S S P T P T C L L
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Q L I M V T N K A I A S Q I S Q I K H F F H C I L V V V C P N S S
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FIG. 14.

v360 v370 v380 v390 v400 v410 v420 v430 v440
SSVGTDVTEGPAHPAPHTRLPFANKEKEPKKKVSELRSELWEKMKLTDIRLEALNSAQDQLRETHNQLEVDLLKAENDRLKVAP
SSVGT:VTE.PAH:..PTRLF:ANKEKEPKKKVSELRSELWEKMKLTDIRLEALNSAQDQLRETHNQLEVDLLKAENDRLKVAP
SSVGTETVTEPAHSVPBTRLPQANKEKEPKKKVSELRSELWEKMKLTDIRLEALNSAQDQLRETHNQLEVDLLKAENDRLKVAP
^10 ^20 ^30 ^40 ^50 ^60 ^70 ^80 ^90

v450 v460 v470 v480 v490 v500 v510 v520 v530
GPSSGCTPGQVPGSSALSSPRRLGLALTESFGPSLADTDLSPMDGISTCGPKKEVTLRVVVRMPPQHIKGDLLKQOEFLGCSKVSGKV
GPSSG.TPGQVPGSSALSSPRRLGLAL:8:F:PSL:DTDLSPMDGISTCG:KEVTLRVVVRMPPQHIKGDLLKQOEFLGCSKVSGKV
GPSSGCTPGQVPGSSALSSPRRLGLALSFPSPSLTDTLSPMDGISTCGSKKEVTLRVVVRMPPQHIKGDLLKQOEFLGCSKVSGKV
^100 ^110 ^120 ^130 ^140 ^150 ^160 ^170 ^180

v540 v550 v560 v570 v580 v590 v600 v610 v620
DWRQLDEAVFQVFKDYISKMDPASTLGLSTESIEGYSISEVTRVLDARPEMPPCRRGVNNISVSLKGLKEKCVDSLVTETLIPKPMHQH
DWRQLDEAVFQVFKDYISKMDPASTLGLSTESIEGYS:SEVTRVLDARPEMPPCRRGVNNISV:LKGLKEKCVDSLVTETLIPKPMHQH
DWRQLDEAVFQVFKDYISKMDPASTLGLSTESIEGYSISEVTRVLDARPEMPPCRRGVNNISVALKGLKEKCVDSLVTETLIPKPMHQH
^190 ^200 ^210 ^220 ^230 ^240 ^250 ^260 ^270

v630 v640 v650 v660 v670 v680 v690 v700 v710
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YISLLKRRRLVLSGPGSGTGKTYLTNRLAAYLVERSGREVT:GIVSTFNMEQQSCDKDLQLYLSNLANQIDRETGIGDVPLVILLDDLSEA
YISLLKRRRLVLSGPGSGTGKTYLTNRLAAYLVERSGREVTGIVSTFNMEQQSCDKDLQLYLSNLANQIDRETGIGDVPLVILLDDLSEA
^280 ^290 ^300 ^310 ^320 ^330 ^340 ^350 ^360

v720 v730 v740 v750 v760 v770 v780 v790 v800
GSISELVNGALTCYKCPYIIGTTNQPVKMTPNHGLELSFRMLTFSNNVEPANGFLVRYLRRLVSESDSDINAKKELLRLVDWVPKLW
GSISELVNGALTCYKCPYIIGTTNQPVKMTPNHGLELSFRMLTFSNNVEPANGFLVRYLRRLVSESDSD:NAKKEILLRLVDWVPKLW
GSISELVNGALTCYKCPYIIGTTNQPVKMTPNHGLELSFRMLTFSNNVEPANGFLVRYLRRLVSESDSDVNAKKEILLRLVDWVPKLW
^370 ^380 ^390 ^400 ^410 ^420 ^430 ^440 ^450

v810 v820 v830 v840 v850 v860 v870 v880 v890
YELHTFLKXSTSDFLIGPCFFLSCPIGIEDFRTWIDLWNNSIIPYLQEGAKDGIKVBGQKAAWEDPVEWVRDTLPWPSAQDQSKLYH
YELHTFLKXSTSDFLIGPCFFLSCPIGIEDFRTWIDLWNNSIIPYLQEGAKDGIKVBGQKAAWEDPVEWVRDTLPWPSAQDQSKLYH
YELHTFLKXSTSDFLIGPCFFLSCPIGIEDFRTWIDLWNNSIIPYLQEGAKDGIKVBGQKAAWEDPVEWVRDTLPWPSAQDQSKLYH
^460 ^470 ^480 ^490 ^500 ^510 ^520 ^530 ^540

v900 v910 v920 v930 v940 v950 v960
LPPFTVGFHSIASPPEDRTVKDSTPSSLDSDPLMALLKLQEAANYIESPDRETILDPNLQATL
LPPP:VGFHS:ASPPEDRTVKDSTP:SLDSDPLMALLKLQEAANYIESPDRETILDPNLQATL
LPPFVGFHSIASPPEDRTVKDSTPSSLDSDPLMALLKLQEAANYIESPDRETILDPNLQATL
^550 ^560 ^570 ^580 ^590 ^600

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FIG. 15.

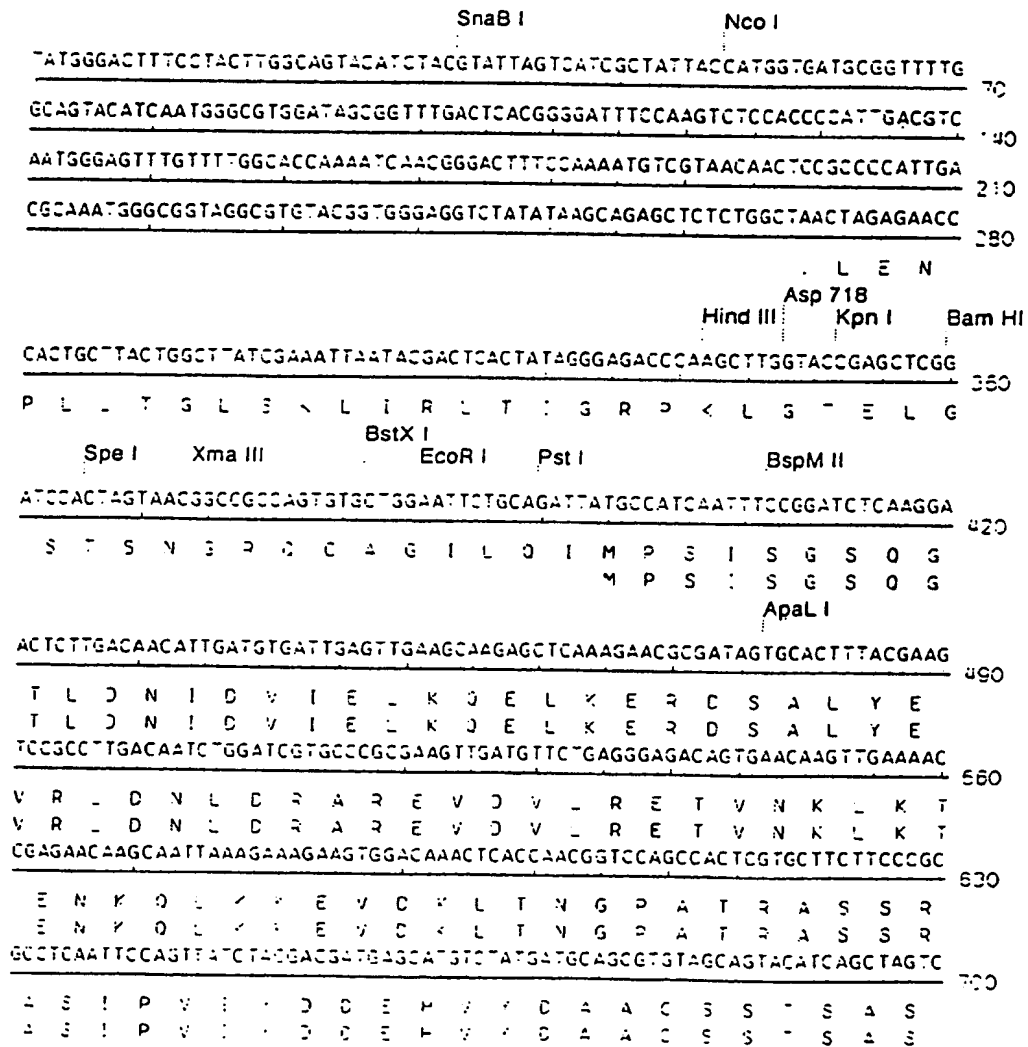


FIG. 15 CONTINUED.

Asu II

AATCTTCGAAACGATCCTCTGGCTSCAACTCAATCAAGTTACTGTAAACSTGGACATCGCTGGAGAAAT 770
 C S S K R S S G C N S I K V T V V V D I A G E :
 C S S K R S S G C N S I K V T V V V D I A G E :

Pvu I

Hpa I

EcoR V

CAGTTCGATCGTTAACCCGACAAAGAGATAATCGTAGGATATCTTGCCATGTCAACCAGTCAGTCATGC 840
 S S I V N P D K E : I V G Y L A M S T S Q S C
 S S I V N P D K E : I V G Y L A M S T S Q S C
 TGGAAAGACATTGATGTTCTATTCTAGGACTATTGAAGTCTACCTATCCAGAATTGATGTGGAGCATC 910

V K D : D V S I L G L F E V Y L S R I D V E H
 V K D : D V S I L G L F E V Y L S R I D V E H

Cla I

Mlu I

AACTTGAATCGATGCTCGTGATTCTATCCTTGGCTATCAAATTGGTGAACCTCGACCGCTCATTTGGAGA 980
 Q L G : D A R D S I L G Y Q I G E L R R V I G D
 Q L G : D A R D S I L G Y Q I G E L R R V I G D
 CTCACAAACCATGATAACCGCCATCCAACTGACATTCTTACTTCCTCAACTACAATCCGAATGTTCTATG 1050

S T T M : T S H P T D : L T S S T T I R M F M
 S T T M : T S H P T D : L T S S T T I R M F M
 CACGCTGCCGCACAGAGTCGCGTAGACAGTCTGGTCTTGATATGCTCTTCCAAAGCAAATGATTCTCC 1120

H G A A Q S R V D S L V L D M L L P K C M I L
 H G A A Q S R V D S L V L D M L L P K C M I L
 AACTCGTCAAGTCAATTTTGACAGAGAGACGTC TGGTGTAGCTGGAGCAACTGGAAATGGAAAGAGCAA 1190

Q L V K S : L T E R R L V L A G A T G I G K S K
 Q L V K S : L T E R R L V L A G A T G I G K S K

Asu II

ACTGGCGAAGACCCCTGGCTGCTTATGTATCTATTCGAACAAATCAATCCGAAGATAGTATGTGAATATC 1260
 L A K T L A A V S I P T N Q S E C S : V N I
 L A K T L A A V S I R T N Q S E C S : V N I

Bsm I

Bgl II

AGCAATTCCTGAAAACAATAAGAGAAGTTCCTTCAAGTGAACGACGCTTGGAAAAGATCTTGAGAAGCA 1330
 S I P E N N K E E L L O V E R R L E : L R S
 S I P E N N K E E L L O V E R R L E : L R S

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FIG. 15 CONTINUED.

Ava III
 Nsi I Xba I

AAGAATCATGCATCGTAATTCAGATAATATCCCAAAGAATCGAATTCATTGTGTGATCCGTTTTTGC 1400
 K E S C I V : L D N I P K N R I A F V V S V F A
 K E S C I V : L D N I P K N R I A F V V S V F A
 EcoR V

AAATGTCCCACTTCAAAACAACGAAGGTCCATTTGTAGTATGCACAGTCAACCGATATCAAAATCCCTGAG 1470
 V V P L G N N E G P F V V C T V N R Y G I P E
 V V P L G N N E G P F V V C T V N R Y G I P E
 CTTCAAAATTCACCAATTTCAAAATGTCAAGTAATGTCGAATCGTCTCGAAGGATTCATCCTACGTTACC 1540
 L Q I H H N F K M S V M S N R L E G F : L R Y
 L Q I H H N F K M S V M S N R L E G F : L R Y
 TCCGACGACGGCGGTAGAGGATGAGTATCGTCTAACTGTACAGATGCCATCAGAGCTCTCAAAATCAT 1610
 L R R R A V E D E Y R L T V D M P S E L F K I :
 L R R R A V E D E Y R L T V D M P S E L F K I :
 EcoR I

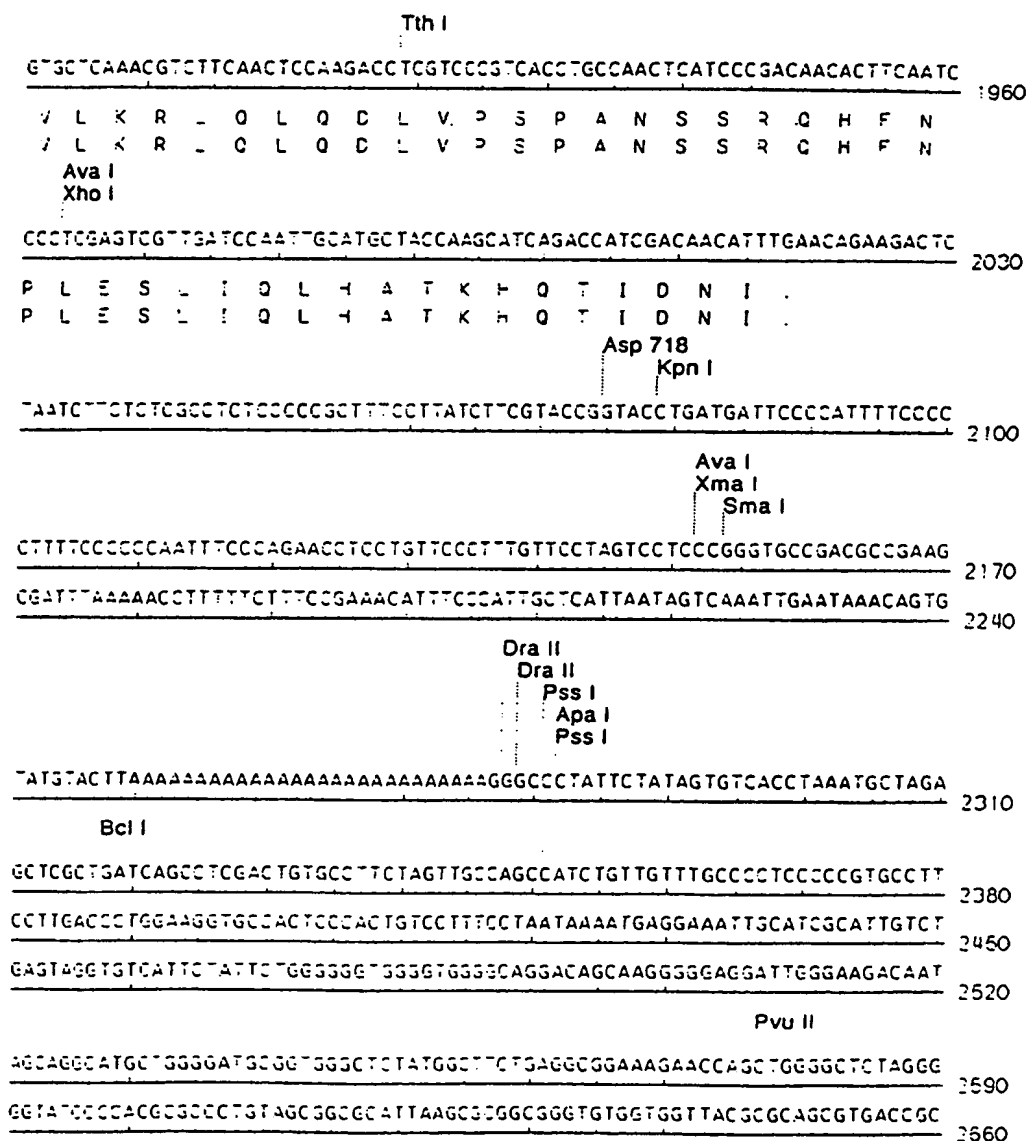
TGACTCTTCCCAATAGCTCTTCAGGCGGTCAATAATTTTATTGAGAAAACGAATTCGTGTGATGAC 1680
 D F F P : A L G A V N N F I E K T N S V D V T
 D F F P : A L G A V N N F I E K T N S V D V T
 Bam HI

GTTSGTCCAAGAGCATGCTTGAACGTGCTCTAACTGTGATGGATCCCGTGAATGGTTCATTCGATTGT 1750
 V G P R A C L N C P L T V D G S R E W F : R L
 V G P R A C L N C P L T V D G S R E W F : R L
 GGAATGAGAACTTCATTCCATATTTGGAACGTGTTGCTAGAGATGGCAAAAAACCTTCGGTCGCTGCAC 1820
 V N E N F : P V L E R V A R D G K K T F G R C T
 V N E N F : P V L E R V A R D G K K T F G R C T
 Bam HI Tth I

TTCTTCGAGGATCCACCGACATCGTCTCTAAAAAATGSCCGTGGTTCGATGGTGA AAAACCCGAGAA 1890
 S F E D P T C : V S K Y W P W F D S E N D E N
 S F E D P T C : V S K Y W P W F D S E N D E N

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FIG. 15 CONTINUED.



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FIG. 15 CONTINUED.

Nae I
TACACTTGCCAGCGCCCTAGCGCCGCTCCTTTGCGTTTCTTCCCTTCCCTTTCTCGCCACGTTGCGCGGC 2730
TTCCCCGTCAAGCTCTAAATCGGGGCATCCCTTTAGGGTTCCGATTAGTGCTTTACGGCACCTCGACC 2800

Dra III
CCAAAAAATTGATTAGGGTGATGGTTACGTTAGTGGGCCATCGCCCTGATAGACGGTTTTCGCCCTTT 2870
GACGTTGGAGTCCCGTTCTTTAATAGTGGACTCTTGTTCCTAACTGGAACAACACTCAACCTATCTCG 2940
GTCTATTCTTTTGATTTATAAGGGATTTTGGGGATTTTCGCCCTATTGGTTAAAAAATGAGCTGATTTAAC 3010
AAAAATTTAAGCGCAATTAATCTGTGAATGTGTGTCAGT"AGGGTG"GGAAAGTCCCCAGGCTCCCCA 3080

Ava III
Nsi I
GGCAGGCAGAGTATGCAAAGCATGCA"CTCAATTAGTCASCAACCAGGTGTGGAAAGTCCCAGGCTCC 3150

Ava III
Nsi I
CCAGCAGGCAGAGTATGCAAAGCATGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGC 3220

Nco I
CCATCCCGCCCTTACTCCGCCCAGTTCGCCCCATTC"CCGCCCATGGCTGACTAATTTTTTTTATTTA 3290

Stu I
Avr II
TGCAGAGGCGGAGGCGCCCTCTGCCTCTGAGCTATTCCAGAAGTAGTGAGGAGGCCTTTTGTGGAGGCTA 3360

Ava I
Xma I
Sma I
Bcl I
GGCTTTTGCAAAAAGCTCCCGGGAGCTTGATATCCATTTTCGGATCTGATCAAGAGACAGGATGAGGAT 3430

Xma III
CGTTTCGATGATTGAACAAGATGGATTGCACGGAGGTTCTCCGCGCGCTTGGGTGGAGAGGCTATTTCGG 3500

Nar I
Bbe I
CTATGACTGGGCACACAGACAACTGCGCTGCTCTGATGCCGCGGTCTCCGCTGTCAGCGCAGGGGCGC 3570

| | |
|-----------------------------------------------------------------------------------------------------------------------------------------|------|
| CCGGTCTTTTSTCAAGACCGACCTGTCCGGTCCCTGGAATGAAGTGCAGGACGAGCGAGCGCGGTAT | 3640 |
| <div style="display: flex; justify-content: space-around;"> <div>Bal I</div> <div>Fsp I</div> <div>Pvu II</div> <div>Tth I</div> </div> | |
| CGTGGCTGGCCACGACGGGCGCTTCTTGCSCAGCTGTGCTCGACGTTGTCACTGAAGCGGGAAGGGACTG | 3710 |
| GCTGCTATTGGGCGAAGTGCCGGGGGAGGATCTCTGTGATCTCACCCTTCTCCGCGAGAAAGTATCC | 3780 |
| ATCATGGCTGATGCAATGCGGCGCTGTCATACGCTTGATCGGGCTACCTGCCCATTCGACCACCAAGCGA | 3850 |
| AACATCGCATCGAGCGAGCAGCTACTCGGATGGAAGCCGGTCTTGTGATCAGGATGATCTGACGAAGA | 3920 |
| BssH II | |
| GCATCAGGGGCTGGGCCAGCGGAACTGTTGCCAGGCTCAAGGGCGGCATGCCCGACGGCGAGGATCTC | 3990 |
| Nco I | |
| GTCGTGACCCATGGCGATGCCCTGTTGCCGAATATCATGCTGGGAAAATGGCGGCTTTCTGGATTCATCG | 4060 |
| <div style="display: flex; justify-content: space-around;"> <div>Nae I</div> <div>Rsr II</div> </div> | |
| ACTGTGGCCCGCTGGGTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGAAGA | 4130 |
| GCTTGGGCGGGAATGGGCTGACCGCTTCTGTGCTTACGGATCGCCGCTCCCGATTCGACGGCGATC | 4200 |
| Asu II | |
| GCCTTCATCGCCTTCTTGACGASTTCTTCTGAGCGGGACTCTGGGGTGGAAATGACCGACCAAGCGAC | 4270 |
| GCCCAACCTGCCATCACGAGATTTCGATTCCACCGCGCCTTCTATGAAAGGTTGGGCTTCGGAATCCTT | 4340 |
| Nae I | |
| TTCCGGGACCGCGGCTGGATGATCTCTCCAGCGCGGGGATCTCATGCTGGAGTTCTTGGCCACCCCAACT | 4410 |
| TGTTATTGAGGCTTATAATGCTTACAAATAAAGCAATAGCATCACAAATTCACAAATAAAGCATTTTT | 4480 |
| <div style="display: flex; justify-content: space-around;"> <div>Bsm I</div> <div>Sal I</div> </div> | |
| TTCACTGCATCTAGTTGTGGTTTGTCCAAACTCATCAATGATCTTATCATGCTGTATACCGTCGACC | 4550 |
| CTAGCTAGAGCTTGGCGTAATCATGGCTATAGGCTTTCTGTGTGAAATTGTTATCGGCTCACAACTC | 4620 |
| CACACAACATACGAGCGCGGAAGCATGAAGTGTAAAGCCCTGGGGTGCTTAATGAGTGAAGCTAACTCACA | 4690 |

FIG. 15 CONTINUED.

Pvu II

AATTGCGTTGCGCTCAGTCTGGGCTTTCCAGTCGGGAAACCTGTCTGCGCAGCTGCATTAAATGAATCGGC 4760
 CAACGCGCGGGGAGAGGCGCTTTTCCGCTATTGGGCGCTCTTCCGCTTCCCTCGCTCACTGACTCGCTCGCT 4830
 CGGTGCTTCGGCTGCGGCGAGCGCTATCAGCTCACTCAAAGCGCGTAATACGGTTATCCACAGAATCAGG 4900
 GGATAACGCAGGAAAGAACAATGTGAGCAAAAGGCCAGCAAAAGGCCAGGAACCGTAAAAAGCGCGCTTG 4970
 CTGGCGTTTTCATAGGCTCGGCGCGCTGACGAGCATCAGAAAAATCGACGCTCAAGTCAGAGGTGGC 5040
 GAAACCCGACAGGACTATGAGATACCAGGCGTTTCCCGCTGGAAGCTCCCTCGTGGCTCTCTCTTCC 5110
 GACCTGCGCGCTTACCGGATACCTGTCCGCGCTTCTCCCTTCGGGAAGCGTGGCGCTTCTCAATGCTCA 5180

ApaL I

CGCTGTAGGTATCTCAGTCTGGTCTAGGTCGTTCCCTCCAAGCTGGGCTGTGTGCACGAACCCCGCTTC 5250
 AGCCCCACCGCTGCGCGCTTATCGGTAACCTATCGTCTTGAGTCCAACCCGGTAAGACACGACTTATCGCC 5320

AlwI

ACTGCGAGCAGCCACTGGTAACAGGATTAGCAGAGCGAGGATATGTAGGCGGTGCTACAGAGTCTTTGAAG 5390
 TGGTGGCTTAACACGGCTACACTAGAAGGACAGTATTTGGTATCTGCGCTCTGCTGAAGCCAGTTACCT 5460
 TCGGAAAAAGAGTTGGTAGCTCTTATCGGCAAAACAAACACCGCTGGTAGCGGTGGTTTTTTGTTTG 5530
 CAAGCAGCAGATTACGCGCAGAAAAAAGGATCTCAAGAAGATCCTTTCATCTTTCTACGGGCTTGAC 5600

BspH I

GCTCAGTGGAAACGAAAACTCAGCTTAAGGGATTTTGGTCATGAGATTATCAAAAAAGGATCTTCACCTAGA 5670
 TCTTTTAAATTAAAAATGAAGTTTAAATCAATCTAAAGTAATATGAGTAACCTTGGTCTGACAGTTA 5740
 CCAATGCTTAATCAGTGAGGCACTATCTCAGCGATCTGTCTATTTGGTTCATCTATAGTTGCCTGACTC 5810
 CCGTCTGTAGATAACTAGATACGGGAGGGCTTACCATCTGGCGCGAGTGGTTCATATGATACCGCGAG 5880
 ACCACGCTCACCGGCTCGAGTTTATCAGCAATAAACCAGCCAGCGGGAAGGAGGAGGCGAGAGAGTGG 5950
 TCTTCAACTTTATCGCGCTCTATGAGTCTATTAATTGTTCGCGGGAAGGCTAGAGTAAAGTAGTTCCCA 6020

FIG. 15 CONTINUED.

Fsp I

GTTAATAGTTTGGSCAACSTTTTSCCAATTGCTACAGGCATCGTGGTGTCACGCTCGTCGTTTGGTATGG 6090
 CTTCAATCAGCTCCGCTTCCCAACGATCAAGGCGAGTTACAATGATCCCCCATGTTTGCACAAAAAGCGT 6160

Pvu I

TAGCTCCTTCGGTCCCTCCGATCGTTTGTCAAGTAAGTTGGCCGCGAGTGTATCACTCATGGTTATGGCA 6230

Sca I

GCACCTGCATAATTCTCTTACTTTCATGCCATCCSTAAGATGCTTTTCTGTGACTGGTGAGTACTCAACCA 6300
 AGTCATTCTGAGAATAGTGTATGGCGCGACCGAGTTGCTCTTGGCCGCGTCAATACGGGATAATACCGC 6370
 GCCACATAGCAGAACTTTAAAGTGTCTCATCATTTGSAACGCTTCTTCGGGGCGAAACTCTCAAGGATC 6440

ApaL I

TTACCGCTGTGAGATCCAGTTGGATGTAACCCACTCGTGCACCCAACTGATCTTCAGCATCTTTTACTT 6510
 TCACCAGCGSTTTCTGGGTGATCAAAAAACAGGAAGGCAAAATGCCGCAAAAAAGGGAATAAGGGCGACACG 6580

Ssp I

BspH I

GAATGTTGAATACTCATACTCTTCTTTTCAATATTATTGAAGCAATTTATCAGGSTTATTGTCTCATG 6650
 AGCGGATACATATTTGAATGTATTTAGAAAAATAACAAATAGGGGTTCGCGGCACATTTCCCGGAAAAG 6720

Sal I

Bgl II

Sal I

TGCCACCTGACGTGACCGATCGGGAGATCTCCCAATCCCTATGGTGGACTCTCAGTACAATCTGCTCT 6790

AlwI

GATGCCGCATAGTTAAGCCAGTATCTGCTCCCTGCTTGTGTGTTGGAGGTGCTGAGTAGTSCGCGAGCA 6860
 AAATTTAAGCTACACACAAGGCAAGGCTTGACCGACAATTCATGAAGAATCTGCTTAGGGTTAGGCGTTT 6930

Nru I

Mlu I

Spe I

TGGCGTGGTTCGCGATGTACGGCCAGATATACCGCTTGACATTTGATTTATTGACTAGTTATTAATAGTAA 7000
 TCAATTCGCGGTCTATTTAGTTATAGCCCATATATGGAGTTCCGCGTTACATAACTTACCGTAAATGSCC 7070
 CGCGTGGCTGACCGGCCAAATGACCCCGGCCATGACGTCATTAATGACGTATGTTCGGTAGTAACGCC 7140

AATAGGGACCTTCCATTTGACGTCAATGGGTGGACTATTTACGGTAAACTGCCCACTTGGCAGTACATCAA 7210

Nde I

GTGTATCATATGCCAAGTACGGCCCTATTGACGTCAATGACGCTAAATGGCCCGCGTGGCATTATGCCC 7280
 AGTACATGACCT

7292

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FIG. 16.

Sst I
TATAAGCAGAGCTCTCTGGCTAACTAGAGAACCCACTGCTTACTGGCTTATCGAAATTAATACGACTCAC 70
Ppa I Hind III Sst I Bam HI Spe I Xma III EcoR I
TATAGGGAGACCCAAAGCTTGTACCGAGCTCGGATCCACTAGTAACGGCCGCCAGTGTGCTGGAATTCTG 140
R P P V C V N S
Bgl II Afl III
CAGATCTTGGCTATCAAATTGGTGAACCTCGACGCGTCATTGGAGACTCCACAACCATGATAACCAAGCCA 210
A D L G Y Q : S E L R R V : G D S T T M : T S H
TCCAACCTGACATTCTTACTTCTTCAACTACAATCCGAATGTTTCATGCACGGTGCCGCACAGAGTCGCSTA 280
P T D : L T S S T T I R M F M H G A A Q S R V
P T D : L T S S T T I R M F M H G A A Q S R V
GACAGTCTGGTCTTGATATGCTTCTTCCAAAGCAAATGATTCTCCAACCTCGTCAAGTCAATTTTGACAG 350
D S L V L D M L L P K Q M : L Q L V K S : L T
D S L V L D M L L P K Q M : L Q L V K S : L T
Bbv II
AGAGACGTCTGGTGTAGCTGGAGCAACTGGAATTGAAAGAGCAAACCTGGCGAAGACCCCTGGCTGCTTA 420
E R R L V L A G A T G I G K S K L A K T L A A Y
E R R L V L A G A T G I G K S K L A K T L A A Y
Asu II Bsm I
TGTATCTATTGGAACAAATCAATCCGAAGATAGTATTGTTAATATCAGCATTCTGAAAACAATAAGAA 490
V S I R T V Q S E D S : V N : S I P E N N K E
V S I R T N Q S E D S : V N : S I P E N N K E
Xmn I Bgl II Ava III
GAATTGCTTCAAGTGGAAACGACGCTTGGAAAAGATCTTGAGAAGCAAAGAATCATGCATCGTAATTCTAG 560
E L L O V E R R L E K I L R S K E S C : V I L
E L L O V E R R L E K I L R S K E S C I V I L
ATAATATCCCAAAGAATCGAATTCGATTGTGTATCCGTTTTSCAAATGTCCCACTTCAAAACAACGA 630
D N I P : N R : A F V V S V F A N V P L Q N N E
D N I P : N R : A F V V S V F A N V P L Q N N E
EcoR V
AGGTCCATTGTAGTATGCACAGTCAACCGATATCAAAATCCCTGAGCTTCAAATTCACCACAATTTCAAA 700
S P F V V C : V N P Y C I P E L O I H N F C
S P F V V C : V N R Y C I P E L O I H N F C

FIG. 16 CONTINUED.

ATGTCAGTAATGTCGAATCGTCTCGAAGGATTTCATCTACGTTACCTCCGACGACGGGCGGTAGAGGATG 770
M S V M S N R L E G F I L R Y L R R R A V E D
M S V M S N R L E G F I L R Y L R R R A V E D
Sst I
AGTATCGTCTAACTGTACAGATGCCATCAGAGCTCTTCAAAATCATTGACTTCTTCCCAATAGCTCTTCA 840
E Y R L T V Q M P S E L F K I I D F F P I A L Q
E Y R L T V Q M P S E L F K I I D F F P I A L Q
EcoR I
GGCCGTCAATAATTTTATTGAGAAAACGAATTCGTGATGTGACAGTTGGTCCAAGAGCATGCTTGAAC 910
A V N N F I E K T N S V D V T V G P R A C L N
A V N N F I E K T N S V D V T V G P R A C L N
Bam HI
TGTCCTCTAACTGTGATGGATCCCGTGAATGGTTCATTGATGTTGGAATGAGAACCTCATTCCATATT 980
C P L T V D G S R E W F I R L V N E N F I P Y
C P L T V D G S R E W F I R L V N E N F I P Y
Afl III Bam HI Tth I
TGGAACGTGTTGCTAGAGATGGCAAAAAACCTTCGGTCTGCACTTCCTTCGAGGATCCACCGACAT 1050
L E R V A R D S K K T F G R C T S F E D P T D I
L E R V A R D S K K T F G R C T S F E D P T D I
Bbv II
CGTCTCTAAAAATGGCCGTGCTTCGATGGTGAAAAACCCGAGAATGTGCTCAAACGTCTTCAACTCCAA 1120
V S K K V P W F D G E N P E N V L K R L Q L Q
V S K K V P W F D G E N P E N V L K R L Q L Q
Tth I Xho I
GACCTCGTCCCGTCACCTGCCAACTCATCCCGACAACACTTCAATCCCTCGAGTCGTTGATCCAATTGC 1190
D L V P S P A N S S R Q H F N P L E S L I Q L
D L V P S P A N S S R Q H F N P L E S L I Q L
Bbv II
ATGCTACCAAGCATCAGACCATCGACAACATTTGAACAGAAGACTCTAATCTTCTCTCGCCTCTCCCCCG 1260
H A T K H O T I D N I
H A T K H O T I D N I
CTTTCCTTATCTTCTACCGTACCTGATGATTCGCCATTTTCCCGCTTTTCCCCCAATTTCCCGAGAAC 1330
Xma I
Sma I
CTCCTGTTCCCTTTTGTCTAGTCTCTCCGSGTGCCGACGCCGAAGCGATTTAAAAACCTTTTCTTTCC 1400
Xmn I
GAAACATTTCCCATTTGCTCATTAATAGTCAAAATTAATAAACAGTGTATGTACTTAAAAAAAAAAAAAAAA 1470

FIG. 16 CONTINUED.

Sst I Bcl I
 AAAAAAAAAAAGGGCCCTATTCATAGTGTCACCTAAATGCTAGAGCTCGCTGATCAGCCTCGACTGTG 1540
 CCTTCTAGTTGCCASCCATCTGTGTTTGGCCCTCCCCCGTGCTTCTTTGACCCGGAAGGTGCCACTC 1610
 CCACTGTCTCTTCTAATAAAATGAGGAAATTGCATCGCATTGTCTGAGTAGGTGTCATTCTATTCTGGG 1660
 Bbv II
 GGGTGGGGTGGSSCAGGACAGCAAGGGGGAGGATTGGGAAGACAATAGCAGGCATGCTGGGGATGC3GTG 1750
 GGCTCTATGGCTTCTGAGSSGGAAGAACCAGCTGGGGCTCTAGGGGGTATCCCCACGCGCCCTGTAGCG 1820
 GCGCATTAAAGCGCGCGGCTGTGGTGGTTACGCGCAGCGTGACCGCTACACTTGCCAGCGCCCTAGCGCC 1890
 CGCTCCTTTTGGCTTTCTTCCCTTCTTTCTCGCCACGTTGCGCGGCTTTCCCGCTCAAGCTCTAAATCGG 1960
 GGCATCCCTTAGGGTTCGATTTAGTGCTTTACGGCACCTCGACCCCAAAAAAC TTGATTAGGGTGTATG 2030
 Dra III
 GTTCACGTAGTGGSCCATGSCCCTGATAGACGGTTTTTTCGCCCTTTGACGTTGGAGTCCACGTTCTTTAA 2100
 TAGTGGACTCTTTTCCCAACGGAACAACACTCAACCCATCTCGGCTATTCTTTTGATTATAAGGG 2170
 Xmn I
 ATTTTGGGGATTTCGGCTATTGGTTAAAAAATGAGCTGATTTAACAAAAATTTAACGCGAATTAATTCT 2240
 GTGGAATGTGTCTCAGTTAGGGTGTGGAAGTCCCCAGGCTCCCCAGGCAGGCAGAAGTATGCAAAGCAT 2310
 Ava III
 Nsi I
 GCATCTCAATTAGTCAGCAACCAGGTGTGSAAGTCCCCAGGC*CCCCAGCAGGCAGAAGTATGCAAAGC 2360
 Ava III
 Nsi I
 ATGCATCTCAATTAGTCAGCAACCATAGTCCCGCCCTAACTCCGCCCATCCCGCCCTAACTCCGCCCA 2450
 GTTCCGCCCTTCTCCGCCCATGGCTGACTAATTTTTTTATTTATGACAGAGGCCGAGGCCGCCCTCTGC 2520
 Stu I
 Xma I
 Sma I
 CTCTGAGCTATTCCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCC TAGGCTTTTGCAAAAAGCTCCCGGGA 2590
 Bcl I
 GCTTGATATCTATTCTCGATCTGATCAAGAGACAGGATGAGGATCGTTTCCCATGATTGAACAAGATG 2660
 Xma III
 GATTGCACGCAGGTTCTCCGGCCGCTTGGSTGGAGAGGCTATTCCGGCTATGACTGGGCACAACAGACAAT 2730
 Nar I
 Bbe I
 CGSCTGCTCTGATGCGCGCTTCTCGGCTGTGAGCGCAGGGGCTCCCGCTTCTTTTGTCAAGACCGAC 2800
 CTGTCCGCTGCGCTGAAATGAATGACAGGACGAGGCAGCGCGCTATCGTGGTGGCCACGACGGCGCTT 2870
 Fsp I
 Tth I
 CTTGCGCAGCTGTGCTGACGTTGTCACTGAAGCGGGAAGGGACTGGCTGCTATTGGCGGAAGTGCGGG 2940
 GCAGGATCTCTGTCACTCACTTGTCTCTCCGAGAAAGTATCCATCATGCTGATGCAATGCGGGCG 3010

FIG. 16 CONTINUED.

CTGCATACGCTTATCCGGCTACCTGCCCATTCGACCACCAAGCGAAACATCGCATCGAGCGAGCACGTA 3080
CTCGGATGGAAGCCGCTTGTCSATCAGGATGATCTGGACGAAGAGCATCAGGGGCTCGCGCCAGCCGA 3150

BssH II

ACTGTTCCGCAAGGCTCAAGGCGCSCATGCCCGACGGCGAGSATCTCGTCGTGACCCATGGCGATGCCTGC 3220

Rsr II

TTGCCGAATAATCATSGTGSAATATGGCCGCTTTTCTGGATTTCATCGACTGTGGCCGGCTGGGTGTGGCGG 3290
ACCGCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGAAGAGCTTGGCGGCGAATGGGCTGACCG 3360
CTTCCTCGTGCTTACGGTATCGCCGCTCCCGATTGGCAGCGCATCGCCTTCTATCGCCTTCTTGACGAG 3430

Asu II

TTCTTCTGAGCGSSACTCTGGGGTTGAAATGACCGACCAAGCGAGCGCCCAACCTGCCATCAGGAGATTT 3500
CGATTCCACCGCCGCTTCTATGAAAGGTTGGGCTTCGGAATCGTTTTCCGGGACGCCGGCTGATGATC 3570
CTCCAGCGCGGGATCTCATGCTGAGTTCTTCCGCCACCCCACTTGTATTGTCAGCTTATAATGGTT 3640

Bsm I

ACAAATAAAGCAATAGCATCACAAATTCACAAATAAAGCATTTTTTCACTGCAATTCAGTTGTGGTTT 3710

Sna I

GTCCAAACTCATCAATGTATCTTATCATGTCTGTATACCGTCCGACCTCTAGCTAGAGCTTGGCGTAATCA 3780
TGGTCATAGCTGTTTCCCTGTGTGAAATTTTATCCGCTCACAAATTCACACAACATACGAGCCGGAAGCA 3850
TAAAGTGAAAGCCTGGGGTGCCCTAATGAGTGAGCTAACTCACATTAATTGCGTTGCGCTCACTGCCCGC 3920

Sbo I

TTTCAGTCCGGAAACCTGTCTGTCCAGCTGCATTAATGAATCGGCCAACGCGCGGGAGAGGCGGTTTG 3990
CGTATTGGGCGCTCTTCCGCTTCCCTCGCTCACTGACTCGCTGCGCTCGGTCGTTCCGGCTGCGGCGAGCGG 4060

AII III

TATCAGCTCACTCAAAGGCGGTAATACGGTTATCCACAGAATCAGGGGATAACGCAGGAAAGAATATGTG 4130
AGCAAAAGGCCAGCAAAAGGCCAGGAACCGTAAAGGCCGCTTGTGGCGTTTTCCATAGGCTCCGC 4200
CCCCCTGACGAGCATCACAAAAATCGACGCTCAAGTCAGAGGTTGGCGAAACCCGACAGGACTATAAAGAT 4270
ACCAGGCGTTTCCCGCTGGAAGCTCCCTCGTGCCCTCTCTGTTCCGACCTTGCCGCTTACCGGATACCT 4340
GTCCGCTTTCTCCCTTCGGGAAGCGTGGCGCTTTCTCAATGCTCACGCTGTAGGTATCTCAGTTGGGTG 4410

ApaL I

TAGGTCGTTCCCTCCAAAGCTGGGCTGTGTGCACGAACCCCGCTTCAGCCCGACCGCTGCGCCTTATCCG 4480

AlwI I

GTAACTATCTCTTGAAGTCCAAACCCGGTAAGACACGACTTATGCCACTGGCAGCAGCCACTGGTAACAG 4550
GATTAGCAGAGCTAGGTATGTAGGCGGTCTACAGAGTTCTTGAAGTGGTGGCTTAACACGGCTACACT 4620
AGAAGGACAGTATTGGTATCTCCGCTCTGCTGAAGCCAGTTACCTTCGGAAGAGAGTTGGTAGCTCTT 4690
GATCCGGCAAAACCAACCAACCGCTGATAGCGGTGTTTCTTCTGCAAGCAGCAGATTACGCGCAGAAA 4760
AAAAGGATCTCAAGAGATCTTATCTTTCTACGGGTCTGACGCTCAGTGGAACGAAAACTCACGT 4830

BspH I

TAAAGGATTTTGGTCAAGATTAACAAAAAGGATCTTCACTAGATCTTTTAAATTAATAAAGAT 4900

FIG. 16 CONTINUED.

TTAAATCAATCTAAAGTATATATGAGTAAACTTGGTCTGACAGTTACCAATGCTTAATCASTGAGGCACC 497C
TATCTCAGCGATCTGTCTATTCTTTCATCCATAGTTGCCTGACTCCCCGTCGTGTAGATAACTACGATA 504C
Ppa I
CGGGAGGGCTTACCATCTGSCCCAGTCTGCAATGATACCGGAGACCCACGCTCACCGGCTCCAGATT 511C
TATCAGCAATAAACCAGCCAGCCGSAAGG3CCGAGCGCAGAAGTGGTCTTGCAACTTTATCCGCTCCAT 518C
Fsp I
CCAGTCTATTAATTGTTGCCGGGAAGCTAGAGTAAGTAGTTGCCAGTTAATAGTTTGGCAACGTTGTT 525C
GCCATTGCTACAGGCATCTGGTGTAC3CTCGTCTTGGTATGGCTTCATTCAGCTCCGGTCCCAAC 532C
Pvu I
GATCAAGGCGAGTTACATGATCCCCAT3TTGTGCAAAAAAGCGTTAGCTCCTTCGGTCTCCGATCGT 539C
TGTGAGAAGTAAGTTGGCCGCGAGTGTATCACTCATGGTTATGGCAGCACTGCATAATTCTCTTACTGTC 546C
Sca I Sbo I
ATGCCATCCGTAAGATGCTTTCTGTGACTGGTGAGTACTCAACCAAGTCATTCTGAGAATAGTGTATGC 553C
GGC3ACCGAGTT3CTCTTGGCCGCGTCAATACGGGATAATACCGCGCCACATAGCAGAACTTTAAAGT 560C
Xmn I
GCTCATCATTGGAAAACGTTCTTCCGGGCGSAAAACCTCTCAAGGATCTTACC3CTGTTGAGATCCAGTTCC 567C
EcoK ApaL I
ATGTAACCCACTCGTGCACCCAACTGATCTTCAGCATCTTTTACTTTACCAGCGTTTCTGGGTGAGCAA 574C
AAACAGGAAGGCAAAATGCCGCAAAAAAGG3AATAAGGGCGACACGGAAATGTTGAATACTCATACTCTT 581C
Ssp I BspH I
CCTTTTCAATATTATTGAAGCATTATCAGGGTTATTGTCTCATGAGCGGATACATATTTGAATGTATT 588C
TAGAAAAATAAACAAATAGGGGTTCCGCGCACATTTCCCGSAAAAGTGCCACCTGACGTCGACGGATCGG 595C
Bgl II AlwN I
GAGATCTCCCGATCCCTATGGTCSACTCTCAGTACAATCTGCTCTGATGCCGCATAGTTAAGCCAGTAT 602C
CTGCTCCCTGCTTGTGTGTTGCAAGTCTGCTGAGTAGTGC3GAGCAAAATTTAAGCTACAACAAGGCAAG 609C
Nru I
GCTTGACCGACAATTGCATGAAGAATCT3CTTAGGGTTAGGCGTTTTTGGCTGCTTCCGATGTACGGGC 616C
Afl III Mlu I Spe I
CAGATATACCGCTGACATTGATTATTGACTAGTTATTAATAGTAATCAATTACGGGGTCATTAGTTTAT 623C
AGCCCATATATGGAGTTC3CGCTTACATACCTACGGTAAATGGCCGCGCTGGCTGACCGCCCAACGACC 630C
CC3CCCATTGAC3TCAATAATGACGTAT3TCCCATAGTAACGCCAATAGGGACTTTCCATTGAC3TCA 637C
Nde I
ATGGTGGACTATTTACGGTAAACTGCC3ACTTGG3AGTACATCAAGTGTATCATATGCCAAGTAC3CC 644C
CCTATTGAC3TCAATGAC3TAAATGGCCGCGCTG3CATTAAGCCAGTACATGACCTTATGG3ACTTTC 651C
SnaB I
CTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGTATGCGGTTTTGGCAGTACATCAA 658C
TGGCGTGGATAGCGGTTTGACTCACGGG3ATTTC3AAGTCTCCACCCCATTTGACGTCAATGGGAGTTTG 665C
TTTTGGCACCAAAATCAACGGGACTTTCC3AAATGTCTGAACAACTC3GCCCATTTGACGCAAAATGGGCG 672C
GTAGGCGTGTACGGTGGGAGGTCTA 674C

FIG. 17.

GGTCCGCAACTTTATCCGCTCCATCCAGTCTATTAATTGTTGCCGGAAGCTAGAGTAAGTAGTTCGC 70
Fsp I
CAGTTAATAGTTTSCGCAACGTTGTTGCCATTGCTACAGGCATCGTGGTGTACGCTCGTCGTTTGGTAT 140
GGCTTCATTTCAGCTCCGGTTCCCAACGATCAAGGCGAGTTACATGATCCCCATGTTGTGCAAAAAGCG 210
Pvu I
GTTAGCTCCTTCGGTCTCCGATCGTTGTGAGAAGTAAGTTGGCCGCGAGTGTATCACTCATGGTTATGG 280
Sca I
CAGCACTGCATAATTCTCTTACTGTCATGCCATCCGTAAGATGCTTTCTGTGACTGGTGAGTACTCAAC 350
CAAGTCATTCTGAGAATAGTGATGCGGCGACCGAGTTGCTCTTGCCCGGCGTCAATACGGGATAATACC 420
GCSCCAGATAGCAGAACTTTAAAAGTGCTCATCATTGGAAAACGTTCTTCGGGGCGAAAACCTCTCAAGGA 490
ApaL I
TCTTACCGCTGTTGAGATCCAGTTCGATGTAACCCACTCGTGACCCCAACTGATCTTCAGCATCTTTTAC 560
TTTACCAGCGTTTCTGGGTGAGCAAAAACAGGAAGGCAAAATGCCSAAAAAAGGSAATAAGGGCGACA 630
Ssp I BspH I
CGGAAATGTTGAATACTCATACTCTTCTTTTCAATATTATTGAAGCATTTATCAGGGTTATTGTCTCA 700
TGAGCGGATACATATTTGAATGTATTAGAAAAATAAACAAATAGGGGTTCGCGCACATTTCCCCGAAA 770
Sal I Bgl II Sal I
AGTGCCACCTGACGTGACGGATCGGGAGATCTCCGATCCCCATGGTGGACTCTCAGTACAATCTGCT 840
AlwI
CTGATGCCGCATAGTTAAGCCAGTATCTGCTCCCTGCTTGTGTTGGAGGTGCGTCTAGTAGTGGCGGAG 910
CAAAA'TTAAGCTACAACAAGGC AAGGCTTACCSACAATTGCATGAAGAATCTGCTTAGGGTAGGCGT 980
Nru I Mlu I Spe I
TTTSCGCTGCTTCGCGATGTACCGGCCAGATATACCGGTTGACATTGATTATTGACTAGTTATTAATAGT 1050
AATCAATTACGGGGTCATTAGTTCATAGCCCATATATGGAGTTCCGCGTTACATAACTACGGTAAATGG 1120

FIG. 17 CONTINUED.

CCCGCTGGCTGACCGCCCAACGACCCCGCCATTGACGTCAATAATGACGTATGTTCCCATAGTAACG 1190
CCAATAGGGACTTTCCATTGACGTCAATGGGTGGACTATTTACGGTAAACTGCCCACTTGGCAGTACATC 1260
Nde I
AAGTGATCATATGCCAAGTACGCCCCCTATTGACGTCAATGACGCTAAATGGCCCGCCTGGCATTATGC 1330
SnaB I Nco I
CCASTACATGACCTTATGGGACTTTCCTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATG 1400
GTGATGCGGTTTTGGCAGTACATCAATGGGCGTGGATAGCGGTTTGACTCACGGGGATTTCCAAGTCTCC 1470
ACCCCATTGACGTCAATGGGAGTTTGTTTTGGCACCAAAAATCAACGGGACTTTCCAAAATGTCTGAACAA 1540
CTCCGCCCCATTGACGCAAAATGGGCGGTAGGCGTGTACGGTGGGAGGTCTATATAAGCAGAGCTCTCTGG 1610
Hind III
CTAACTAGAGAACCCTACTGCTTACTGGCTTATCGAAATTAATACGACTCACTATAGGGAGACCCCAAGCTT 1680
L E N D L L T G L S K L I R L T I G R P K L
Asp 718 Kpn I Bam HI Spe I Xma III BstX I EcoR I Pst I
GGTACCGAGCTCGGATCCACTAGTAACGGGCGCCAGTGTGCTGGAATCTGTCAGATCGCCGCTGCCACCT 1750
G T E L S S T S N G R Q C A G I L Q I A A A T
Ava I
Xma I
Sma I
Pvu II
CAACCTTCGGACAACATTCGCTAAGATCCCGGGATACTCATCTATTCTCCACACTTATCAGTCTCAGC 1820
S T F G Q H S L R S P G Y S S Y S P H L S V S A
Spe I
Sal I
TGATAAGGACACAATGTCTATGCACTCACAGACTAGTCCGACGACCTTCTTCACAAAAACCAAGCTATTCA 1890
D Y D T M S M F S C T S R R P S S Q K P S Y S
M E M F S Q T S R R P S S C K P S Y S
GGCCAAATTCATTACCTTGAATGCTAAATGCCACCTTCAAGAGTTTCACTCCACCGAGCACAGAAATGGCGG 1960
S Q F H S L C R K C H L Q E F T S T E F R M A
S Q F H S L C R K C H L Q E F T S T E F R M A

FIG. 17 CONTINUED.

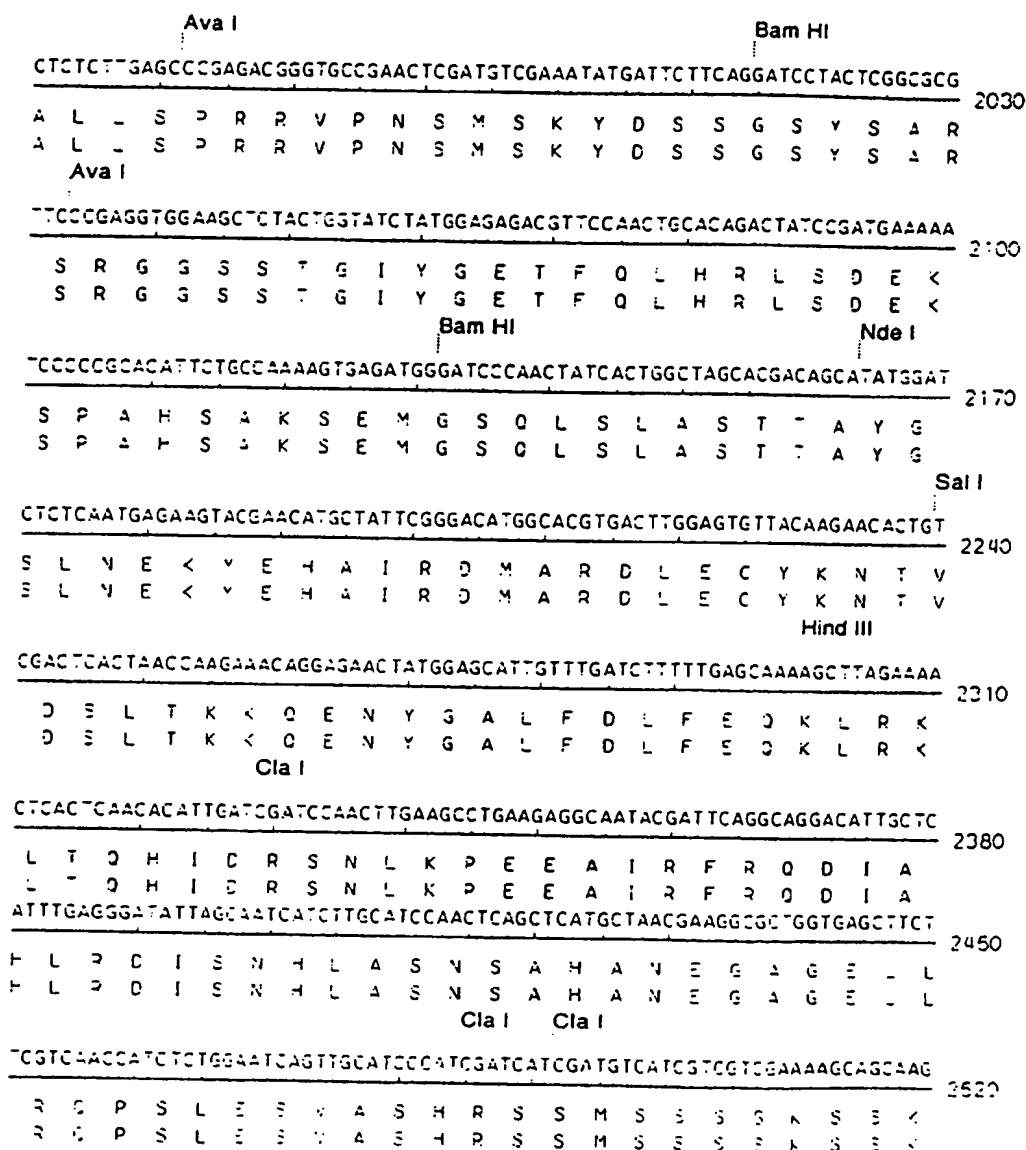


FIG. 17 CONTINUED.

Bam HI

CAGGAGAAGATCAGCTTGAGCTCGTTTGGCAAGAACAAGAAGAGCTGGATCCGCTCCTCACTCTCCAAGT 2590

Q E K I S L S S F G K N K K S W I R S S L S K
Q E K I S L S S F G K N K K S W I R S S L S K

Nde I BspM II

TCACCAAGAAGAAGAACAAGAACTACGACGAAGCACATATGCCATCAATTTCCGGATCTCAAGGAAGCTCT 2660

F T K K K N K N Y D E A H M P S I S G S G G T L
F T K K K N K N Y D E A H M P S I S G S G G T L

ApaL I

TGACAACATTGATGTGATTGAGTTGAAGCAAGAGCTCAAAGAACGCGATAGTGCACTTTACGAAGTCCGC 2720

D N : D V I E L K Q E L K E R D S A L Y E V R
D N : D V I E L K Q E L K E R D S A L Y E V R

CTTGACAATCTGGATCGTGCCCGCAAGTTGATGTTCTGAGGGAGACAGTGAACAAGTTGAAAACCSAGA 2800

L D N L D R A R E V D V L R E T V N K L K T E
L D N L D R A R E V D V L R E T V N K L K T E

ACAAGCAATTAAAGAAAGAAGTGGACAACTCACCAACGGTCCAGCCACTCGTGCTTCTTCCCGCGCCTC 2870

N K Q L < K E V D K L T N G P A T R A S S R A S
N K Q L < K E V D K L T N G P A T R A S S R A S

AATTCAGTTATCTACGACGATGAGCATGTCTATGATGCAACGCTAGCATACATCAGCTAGTCAATCT 2940

I P V I Y D D E H V Y D A A C S S T S A S Q S
I P V I Y D D E H V Y D A A C S S T S A S Q S

Asu II

TCGAAACGATCCTCTGGCTGCAACTCAATCAAGGTTACTGTAAACGTGGACATCGCTGGAGAAATCAGTT 3010

S K R S S G C N S I K V T V N V D I A G E I S
S K R S S G C N S I K V T V N V D I A G E I S

Pvu I Hpa I EcoR V

CGATCGTTAACCCTGACAAAGAGATAATCGTAGGATATCTTGCCATGTCACCCAGTCAGTCAATGCTGGAA 3080

S : I N P D K E : I V G Y L A M S T S D S C L K
S : I N P D K E : I V G Y L A M S T S D S C L K

AGACATTGATGTTCTATTCTTAGGACTATTTGAAGTCTACCTATCCAGAAATTGATGTGGAGCATCAACT 3150

D : D V S I L S L F E V Y L S R : D V E H C L
D : D V S I L S L F E V Y L S R : D V E H C L

Cla I
 GGAATCGATGCTCGTGATTCTATCCTTGGCTATCAAATTGGTGAACCTCGACGCGTCATTGGAGACTCCA
 3220
 G I D A R D S : L G Y O I G E L R R V I G D S
 G I D A R D S : L G Y Q I G E L R R V : G D S
 CAACCATGATAACCCAGCCATCCAACCTGACATTCTTACTTCTCAACTACAATCCGAATGTTTCATGCACGG
 3290
 T T M : T S H P T D I L T S S T T I R M F M H G
 T T M : T S H P T D I L T S S T T I R M F M H G
 TGGCGCACAGAGTCCGCTAGACASTCTGGTCCTTGATATGCTTCTTCAAAGCAAATGATTCTCCAACCTC
 3360
 A A Q S R V D S L V L D M L L P K Q M I L Q L
 A A Q S R V D S L V L D M L L P K Q M I L Q L
 GTCAAGTCAATTTTGACAGAGAGACGCTGGTGTTAGCTGGAGCAACTGGAATTTGAAAGAGCAAACCTGG
 3430
 V K S : L T E R R L V L A G A T G : G K S K L
 V K S : L T E R R L V L A G A T G : G K S K L
 Asu II
 Bsm I
 CGAAGACCCTGGCTGCTTATGTATCTATTCGAACAAATCAATCCGAAGATAGTATTGTTAATATCAGCAT
 3500
 A K T L A A Y V S I R T N Q S E D S : V N I S :
 A K T L A A Y V S I R T N Q S E D S : V N I S :
 Bgl II
 CCTGAAAACAATAAAGAAGAATTGCTTCAAGTGGAACGACGCCCTGGAAAAGATCTTGAGAAGCAAAGAA
 3570
 P E N N K E E L L O V E R R L E K I L R S K E
 P E N N K E E L L O V E R R L E K I L R S K E
 Ava III
 Nsi I
 Xba I
 CATGCATCGTAATTCTAGATAATATCCCAAAGAATCGAATTGCATTGTTGTATCCGTTTTTGCAAATG
 3640
 S C I V I L C V : P K N R I A F V V S V F A N
 S C I V I L C V : P K N R I A F V V S V F A N
 EcoR V
 CCACTTCAAAACAACGAAGGTCATTTGTAGTATGCACAGTCAACCGATATCAAATCCCTGAGCTTCA
 3710
 P L O V N E S P F V V C T V N R Y C I P E L Q
 P L O V N E S P F V V C T V N R Y C I P E L Q
 ATTCAACCAATTTCAAATGTCAGTAATGTCGAATCGTCTCGAAGGATTCATCTACGTTACCTCCGA
 3780
 I H H V F K M S V M S N R L E G F I L R Y L R
 I H H V F K M S V M S N R L E G F I L R Y L R

FIG. 17 CONTINUED

CGACGGGCGGTAGAGGATGAGTATCGTCTAACTGTACAGATGCCATCAGAGCTCTTCAAAATCATTGACT 3850
R R A V E D E Y R L T V Q M P S E L F K I I D
R R A V E D E Y R L T V Q M P S E L F K I I D
EcoR I
TCTTCCAATAGCTCTTCAGCCCTCAATAATTTATTGAGAAAACGAATTCTGTTGATGTGACAGTTGG 3920
F F P : A L C A V N N F I E K T N S V D V T V G
F F P : A L C A V N N F I E K T N S V D V T V G
Bam HI
TCCAAGAGCATGCTTGAACGTGCTCTAACTGTCSATGGATCCCGTGAATGGTTCATTCGATTGTGGAAT 3990
P R A C L V C P L T V D G S R E V F I R L V N
P R A C L V C P L T V D G S R E V F I R L V N
GAGAACTTCATTCCATATTTGGAACGTGTTGCTAGAGATGGCAAAAAACCTTCGGTCGCTGCACTTCTC 4060
E N F : P Y L E R V A R D G K K T F G R C T S
E N F : P Y L E R V A R D G K K T F G R C T S
Bam HI Tth I
TCGAGGATCCACCGACATCGTCTCTAAAAAATGGCCGTGGTTCGATGGTGAAAACCCGGAGAATGTGCT 4130
F E D P T D : V S K K W P W F D G E N P E N V L
F E D P T D : V S K K W P W F D G E N P E N V L
Tth I Ava I Xho I
CAAACGTCTTCAACTCCAAGACCTCGTCCCGTCACCTGCCAACTCATCCCGACAACACTTCAATCCCTC 4200
K R L Q L Q D L V P S P A N S S R Q H F N P L
K R L Q L Q D L V P S P A N S S R Q H F N P L
GAGTCGTTGATCCAATTGCATGCTACCAAGCATCAGACCATCGACAACATTTGAACAGAAGACTCTAATC 4270
E S L : Q L H A T K H O T I D N I
E S L : Q L H A T K H O T I D N I
Asp 718 Kpn I
TTCTCTCGGCTCTCCCCCGCTTCTCTATCTTCGTACCGGTACCTGATGATTCCCATTTTCCCCCTTT 4340
Ava I Xma I Sma I
CCCCCAATTCCGAGAACCCTGTTCCTTTGTTCCTAGTCTCCCGGGTGCCGACGCCGAAGCGATT 4410

FIG. 17 CONTINUED.

TAAAAACCTTTTCTTTCCGAAACATTTCCCATTGCTCATTAAAGTCAGTAATAAACAGGTGATGT

Dra II
Dra II
Pss I
Apa I
Pss I

ACTTAAAAAAAAAAAAAAAAAAAAAAAAAAGGGCCCTATTCTATAGTGTCACCTAAATGCTAGAGCTCG

Bcl I

CTGATCAGCCTCGACTGTGCCCTTAGTTCGCAGCATCTGTTGTTTGCCCCGCCCGTGCCTTCCTTG

ACCTTGAAGGTGCCACTGCCACTGTCTTTCTTAATAAAATGAGGAAAATGCATCGCATGTCTGAGTA

EGTGTCAATCTATTCTGGGGGTTGGGTGGGGCAGGACAGCAAGSGGGAGGATTGGGAAGACAATAGCAG

Pvu II

GCATGCTGGGATGGGGTGGGTCTATGGCTCTGAGGCGGAAAGAACCAGCTGGGGGCTCTAGGGGGTAT

CCCCACGCGCCCTGTAGCAGGCATTAAAGCGGGCGGGTGTGGTGGTTACGCGCAGCGTGACCGCTACAC

Nae I

TGCCAGCGCCCTAGCGCCCGCTGCTTTGCGTTTTCTTCCCTTCTTTCTCGCCACGTTGCGCGGCTTTCC

CCGTCAAGCTCAAATCGGGCATCCCTTAGGGTTCCGATTTAGTGCTTTACGGCACCTCGACCCCAAA

Dra III

AAACTTGATTAGGGTGATGGTTCACGTAGTGGGCATCGCCCTGATAGACGGTTTTTCGCCCTTTGACGT

TGGAGTCCACGTCTTTAATAGTGGACTCTTGTTCGAAACTGGAACAACACTCAACCCATCTCGGTC TA

TTCTTTTGATTTATAAGGSATTTGGGGATTTGGGCTATTGGTTAAAAATGAGCTGATTTAACAAAAA

TTAACGCGAATTAATTCGTGGAATGTGTGTCAGTATGGGTGTGGAAAGTCCCCAGGCTCCCCAGGCAG

Ava III
Nsi I

GCAGAAGTATGCAAGCATGCATCTCAATTAGTCAGCAACCAGGTGTGGAAGTCCCCAGGCTCCCCAGC

Ava III
Nsi I

AGGCAGAAGTATGCAAGCATGCATCTCAATTAGTCAGCAACCAGGTGTGGAAGTCCCCAGGCTCCCCAGC

FIG. 17 CONTINUED.

CCGCCCCAACTCCGCCAGTTCGCCCATTTCTCGCCCCATGGCTGACTAATTTTTTTATTTATGCAG 5530
Nco I
AGGCCGAGGCGCCTCTGCCTCTGAGCTATTCCAGAAGTATGAGGAGGCTTTTGGAGGCTAGGCTT 5600
Stu I
Avr II
TTGCAAAAAGCTCCCGGAGCTTGTATATCCATTTTCGGATCTGATCAAGAGACAGGATGAGGATCGTTT 5670
Ava I
Xma I
Sma I
Bcl I
CGCATGATTGAACAAGATGGATTGCACGCAGGTTCTCCGCGCTTSGGTGGAGAGGCTATTTCGGCTATG 5740
Xma III
ACTGCGCACAACAGACAATCGGCTGCTCTGATGCCGCGCTGTTCCGGCTGTCAGCGCAGGGGCGCCCGT 5810
Nar I
Bbe I
TCTTTTGTCAAGACCGACCTGTCCGGTGGCTTGAATGAACGACAGGACGAGGCGCGGCTATCGTGG 5880
Pst I
CTGGCCACGACGCGCTTCCCTTGGCAGCTGTGCTGACGTTGTCACTGAAGCGGGAAGGGACTGGCTGC 5950
Bal I
Fsp I
Pvu II
Tth I
TATTGGGCGAAGTGCCGGGGCAGGATCTCCTGTCTCTCACCTTGCTCTGCCGAGAAAGTATCCATCAT 6020
GGCTGATGCAATGCGGCGGCTGCATACGCTTGATCCGGCTACCTGCCCATTCGACCACCAAGCGAAACAT 6090
CGCATCGAGCGAGCACGTACTCGGATGGAAGCCGGTCTTGTGATCAGGATGATCTGGACGAAGAGCATC 6160
BssH II
AGGGGCTCGCGCCAGCCGAACGTTTGCCAGGCTCAAGGCGCGCATGCCCGACGGCGAGGATCTCGTGGT 6230
Nco I
GACCCATGGCGATGCTTGGTGGCAATATCATGTTGGAAAATGGCGGCTTTTCTGGAATCATCGACTG 6300

Nae I Rsr II

GGCCGGCTGGGTGTGGCGGACCGCTATCAGGACATAGCGTTGGCTACCCGTGATATTGCTGAAGAGCTTG 6370
GCGCCAATGGGCTGACCCTTCTCTGCTGCTTACGGTATCGCCGCTCCCATTCCAGCGCATCGCTT 6440

Asu II

CTATCGCCTTCTTGACGAGTTCTTCTGAGCGGGACTCTGGGGTTCGAAATGACCGACCAAGCGACGCCCA 6510
ACCTGCCATCACGAGATTTCGATTCCACCSCCGCCTTCTATGAAAGGTTGGGCTTCGGAATCGTTTTCCG 6580

Nae I

GGACGCCGGCTGGATGATCTCCAGCGCGGGATCTCATGCTGGAGTTCTTCGCCACCCCAACTTGTTT 6650
ATTGCGAGCTTATAATGGTTACAAATAAAGCAATAGCATCACAAATTCACAAATAAAGCATTTTTTTCAC 6720

Bsm I Sal I

TGCATTCTAGTTGTGGTTGTCCAAACTCATCAATGTATCTTATCATGTCTGTATACCGTCGACCTCTAG 6790
CTAGAGCTTGGCGTAATCATGGTCATAGCTSTTTCCTGTGTGAAATTGTATCCGCTCACAATTCACAC 6860
AACATACGAGCCGGAAGCATAAAGTGTAAGCCTGGGGTGCTTAATGAGTGAGCTAACTACATTAAATTG 6930

Pvu II

CGTTGCGCTCACTGCCCGCTTTCAGTCSGSAAACCTGTCTGCCAGCTGCATTAATGAATCGGCCAACG 7000
CGCGGGGAGAGGCGGTTTSGCGATTSGGCGCTCTTCCGCTTCTCTCGCTCACTGACTCGCTGCGCTCGGTC 7070
GTTCCGCTGCGGCSAGCGGTATCAGCTCACTCAAAGGCGGTAATACGGTTATCCACAGAATCAGGGGATA 7140
ACGCAGGAAAGAACATGTGAGCAAAAGGCCAGCAAAAGGCCAGGAACCGTAAAAAGSCCGCTTGCTGGC 7210
GTTTTCCATAGGCTCCGCCCCCCCTGACGAGCATCACAAAATCGACGCTCAAGTCAGAGGTGGCGAAAC 7280
CCGACAGGACTATAAAGATACCAGGCGTTTCCCCCTGGAAGCTCCCTCGTGGCGCTCTCTGTTCGACCC 7350
TGGCGCTTACCGGATACCTGTCCGCTTTCTCCCTTCGGGAAGCGTGGCGCTTTCTCAATGCTCACGCTG 7420

ApaI

TAGGTATCTCAGTTGCGGTAGGTCTTCCGTCGAAGCTGGGCTGTGTGACGAACCCCCCTCAGCCC 7490
GACCGCTGCGCCTTATCGGTAACATCTGCTTTSAGTCCAACCCGGTAAGACACGACTTATCGCCACTGG 7560

FIG. 17 CONTINUED.

AlwI

CASCAGCCACTGSTAACAGGATTAGCAGAGCGAGGTATGTAGGCGGTGCTACAGAGTCTTGAAGTGGTG 7620
GCCTAAC TACGGCTACACTAGAAAGGACAGTATTTGGTATCTGCGCTCTGCTGAAGCCAGTTACCTTCGGA 7700
AAAAGAGTTGGTASCTCTTGAATCCGCAAAACAAACCACCGCTGGTASCGGTGGTTTTTTGTTTGCAAGC 7770
AGCAGATTACGCGCAGAAAAAAAGGATCTCAAGAAGATCCTTTGATCTTTTCTACGGGGTCTGACGCTCA 7840

BspH I

GTGGAACGAAAACTCACGTTAAGGGATTTTGGTCATGAGATTATCAAAAAGGATCTTCACCTAGATCCTT 7910
TTAAATTAAAAATGAAGTTTAAATCAATCTAAAGTATATATGAGTAAACTTGGTCTGACAGTTACCAAT 7980
GCTTAATCAGTGAGGCACCTATCTCAGCGATCTGTCTATTTGCTCATCCATAGTTGCCCTGACTCCCGT 8050
CGTGTAGATAAC TACGATACGGGAGGGCTTACCATCTGGCCCCAGTGCTGCAATGATACCGCGAGACCCA 8120
CGCTCACCGGCTCCAGATTATCAGCAATAAACCAGCCAGCCGGAAGGGCCGAGCGCAGAAGT 8183



Figure 18 : Phase contrast images of MCF-7 cells transfected with pCB201 (upper) compared to mock (control) transfected MCF-7 cells (bottom).

The control cells are spread out on the tissue culture plastic and exhibit few filopodia outgrowths. The transfected cells appear smaller because they are slightly rounded up and have multiple filopodia outgrowths (arrowhead) per cell.

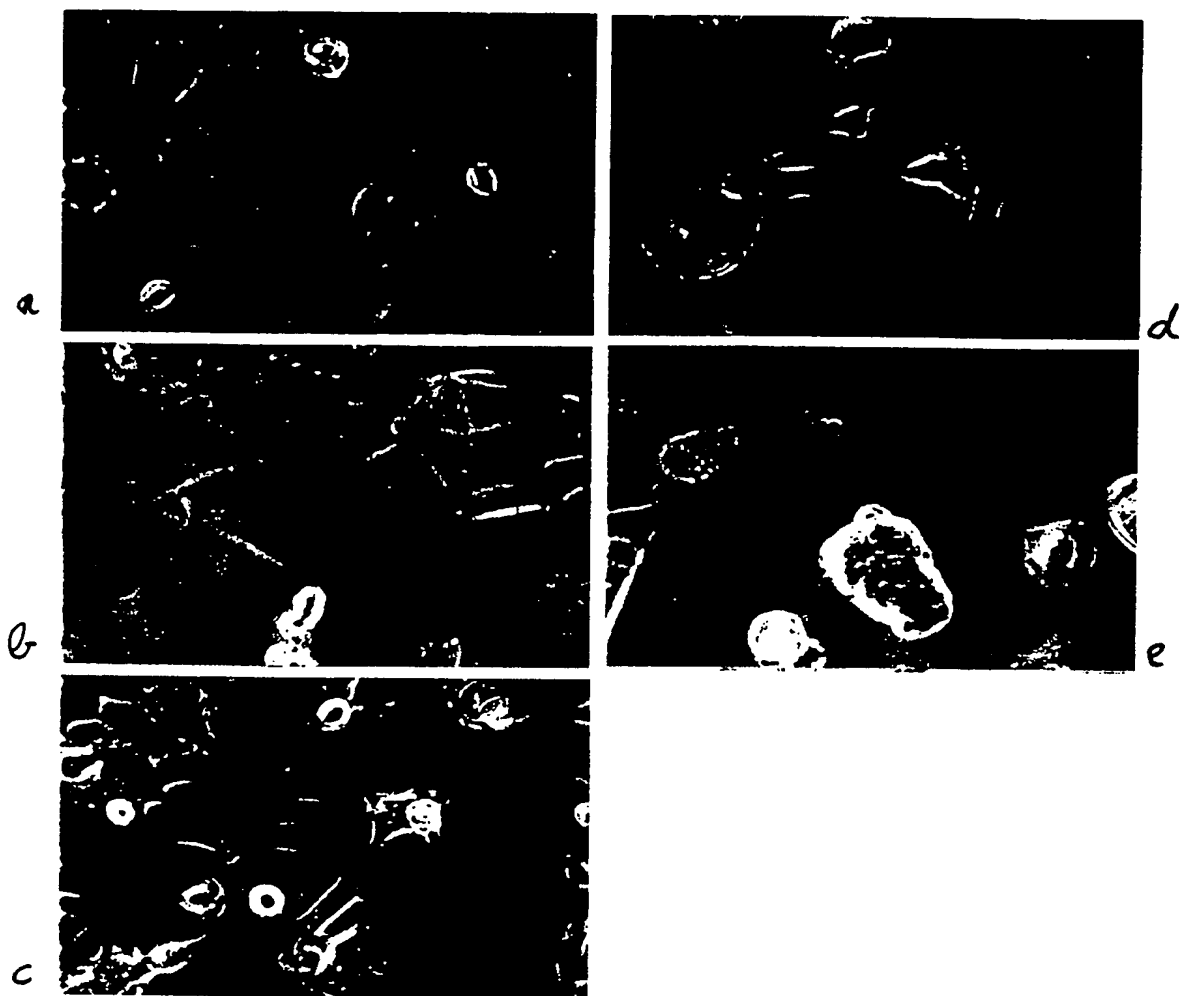


Figure 19 : Phase contrast image of MCF-7 cells, transfected with pcDNA3 (19a), pCDU4 (19b), pCDU3 (19c) pCDU2 (19d) and pTB72 (19e).

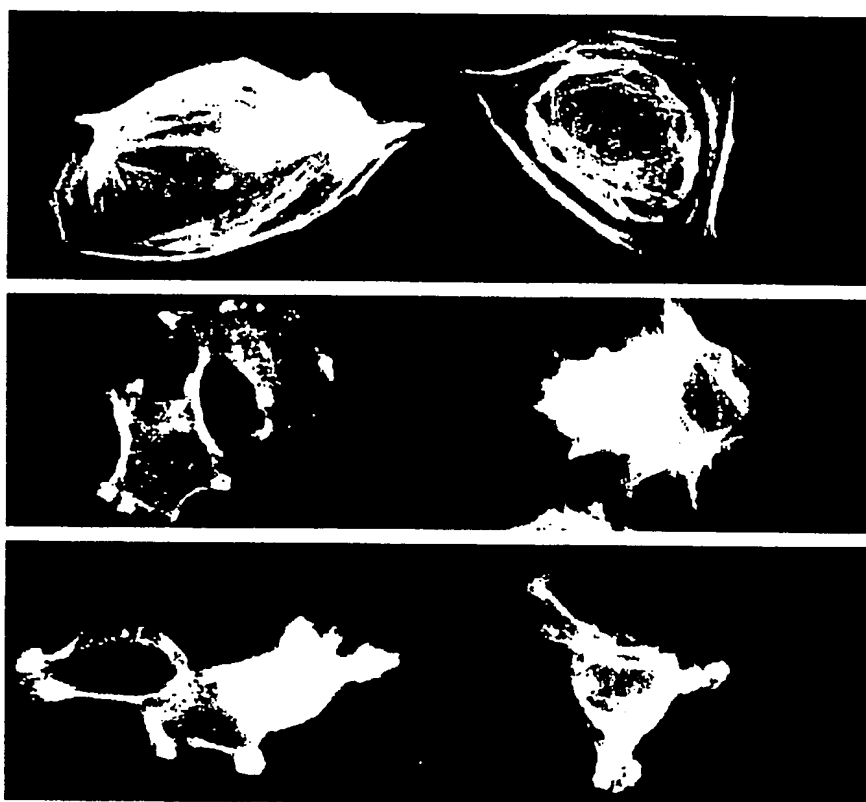


Figure 20 : F- actin pattern (visualized with TRITC-Phalloidin) of MCF-7 cells transfected with pcDNA3.LacZ (top panel) and with pCB201 (middle and lower panel).

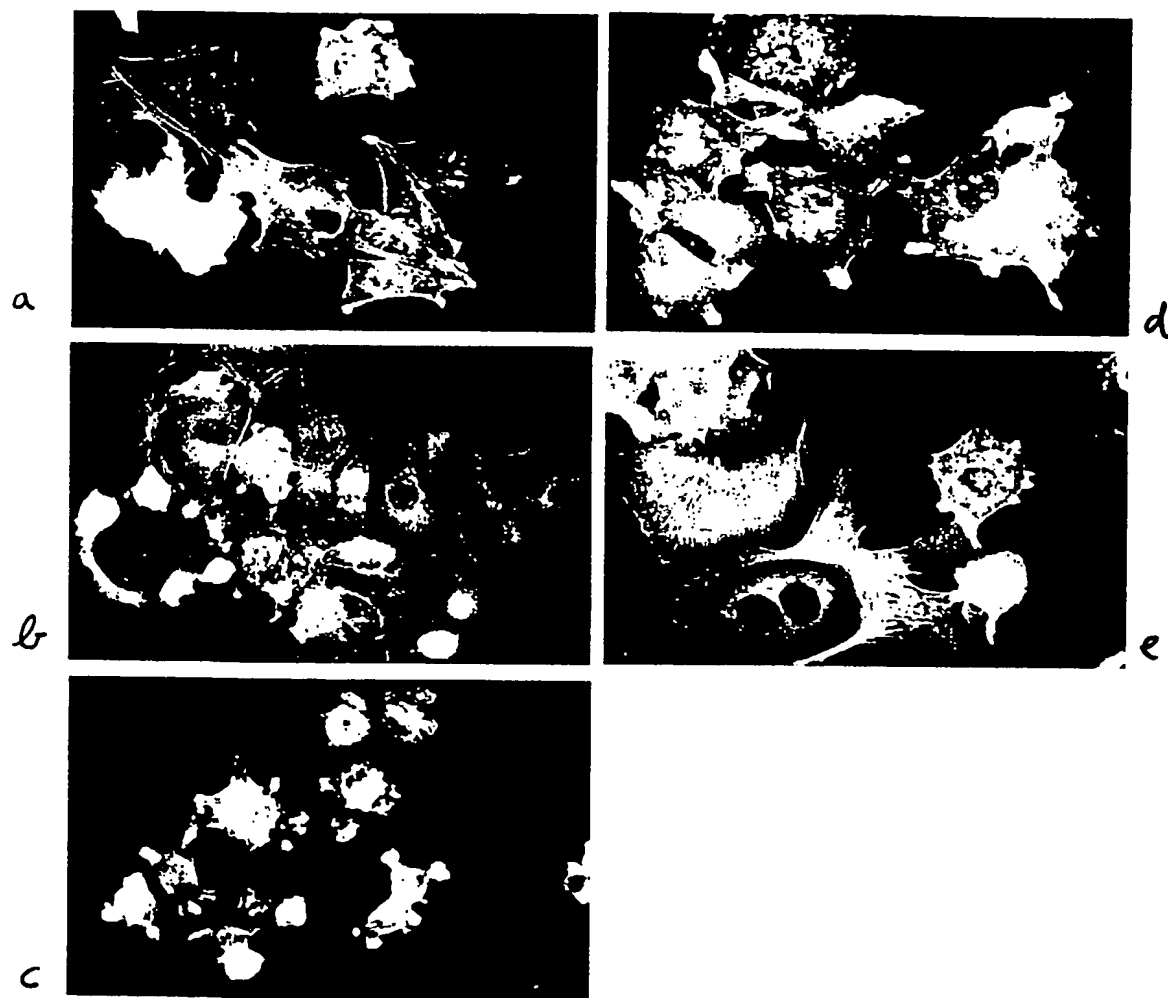


Figure 21 : F- actin pattern Phalloidin (visualized with TRITC-Phalloidin) of MCF-7 cells transfected with pcDNA3 (21a), pCDU4 (21b), pCDU3 (21c) pCDU2 (21d) and pTB72 (21e)

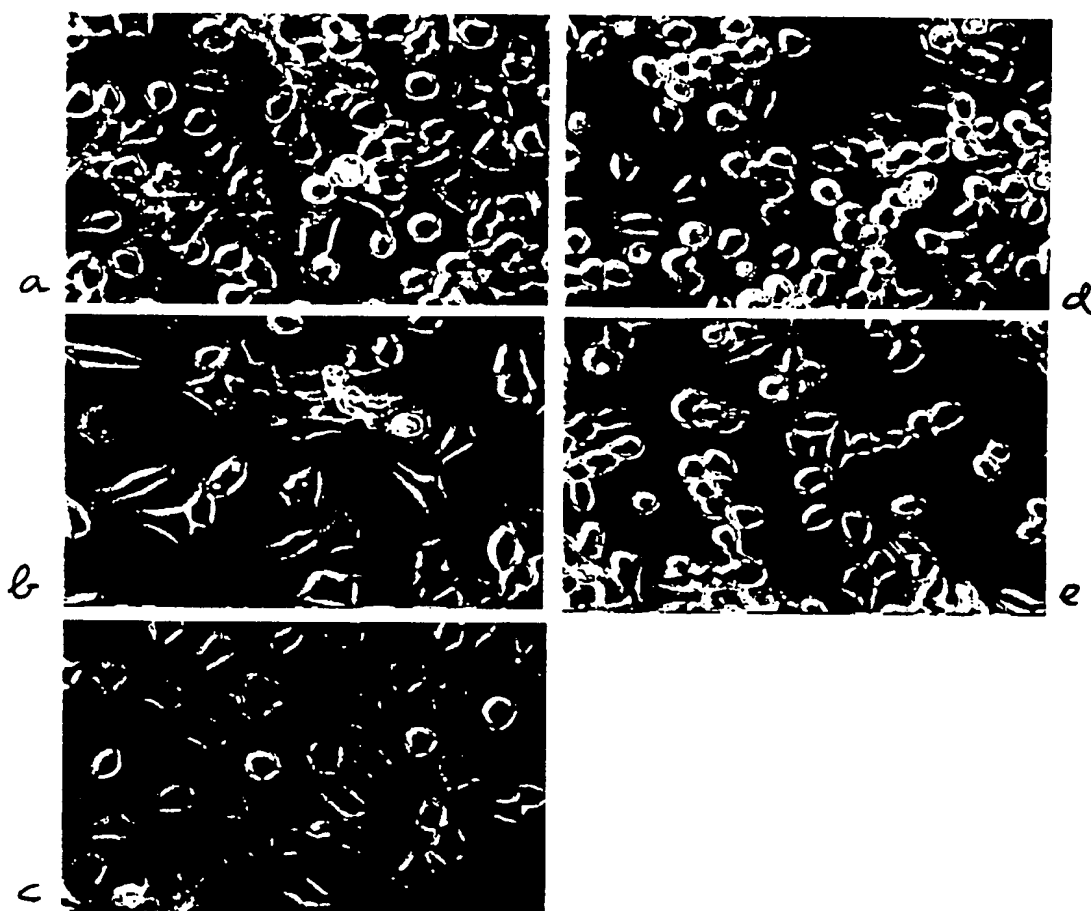


Figure 22 : Phase contrast image of N4 neuroblastoma cells, transfected with pcDNA3 (22a), pCDU4 (22b), pCDU3 (22c) pCDU2 (22d) and pTB72 (22e)

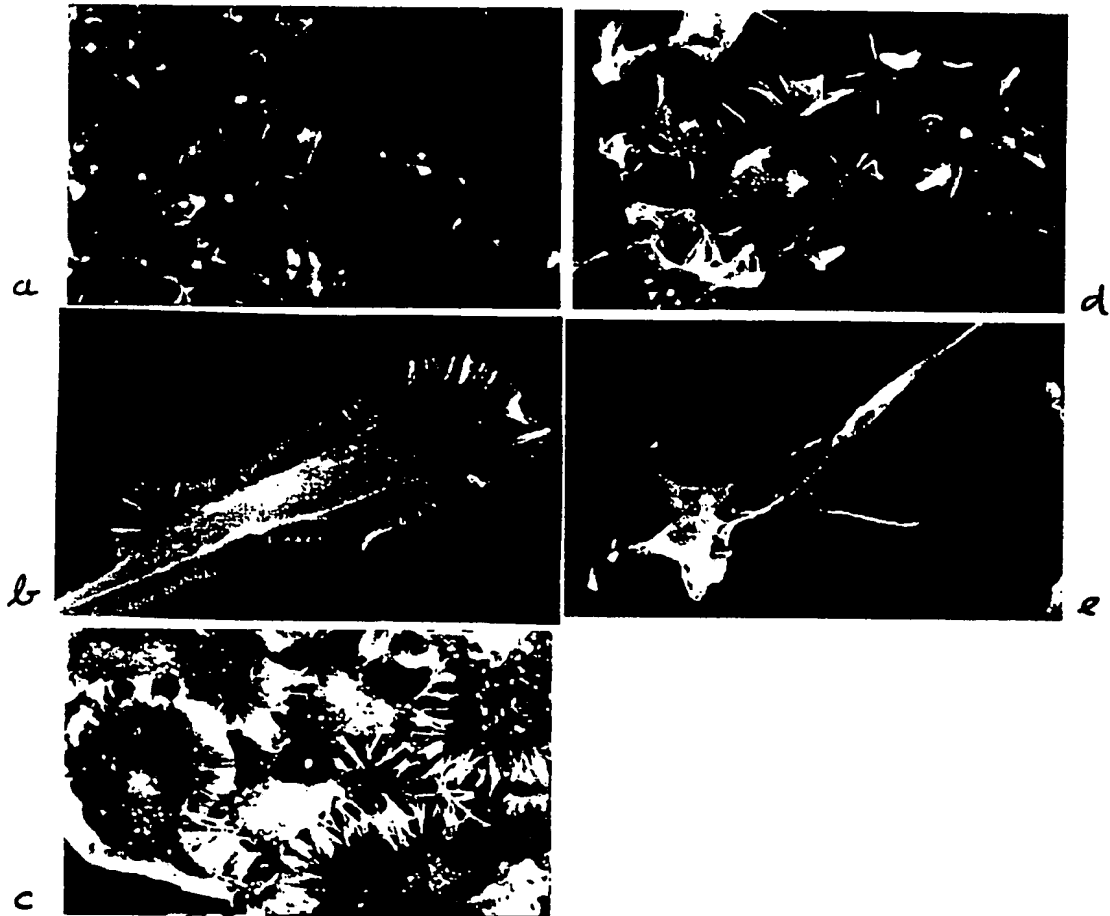


Figure 23 : F- actin pattern Phalloidin (visualized with TRITC-Phalloidin) of N4 neuroblastoma cells transfected with pcDNA3 (23a), pCDU4 (23b), pCDU3 (23c) pCDU2 (23d) and pTB72 (23e)

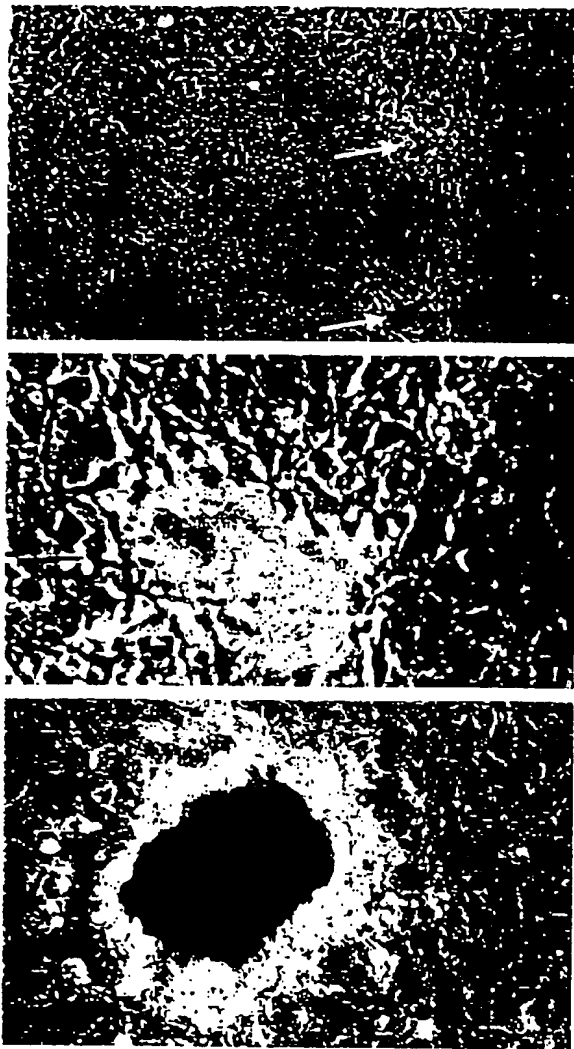


Figure 24 : Phase contrast images of small, medium sized and large foci induced in a monolayer of NIH-3T3 cells by transfection with pCB201.

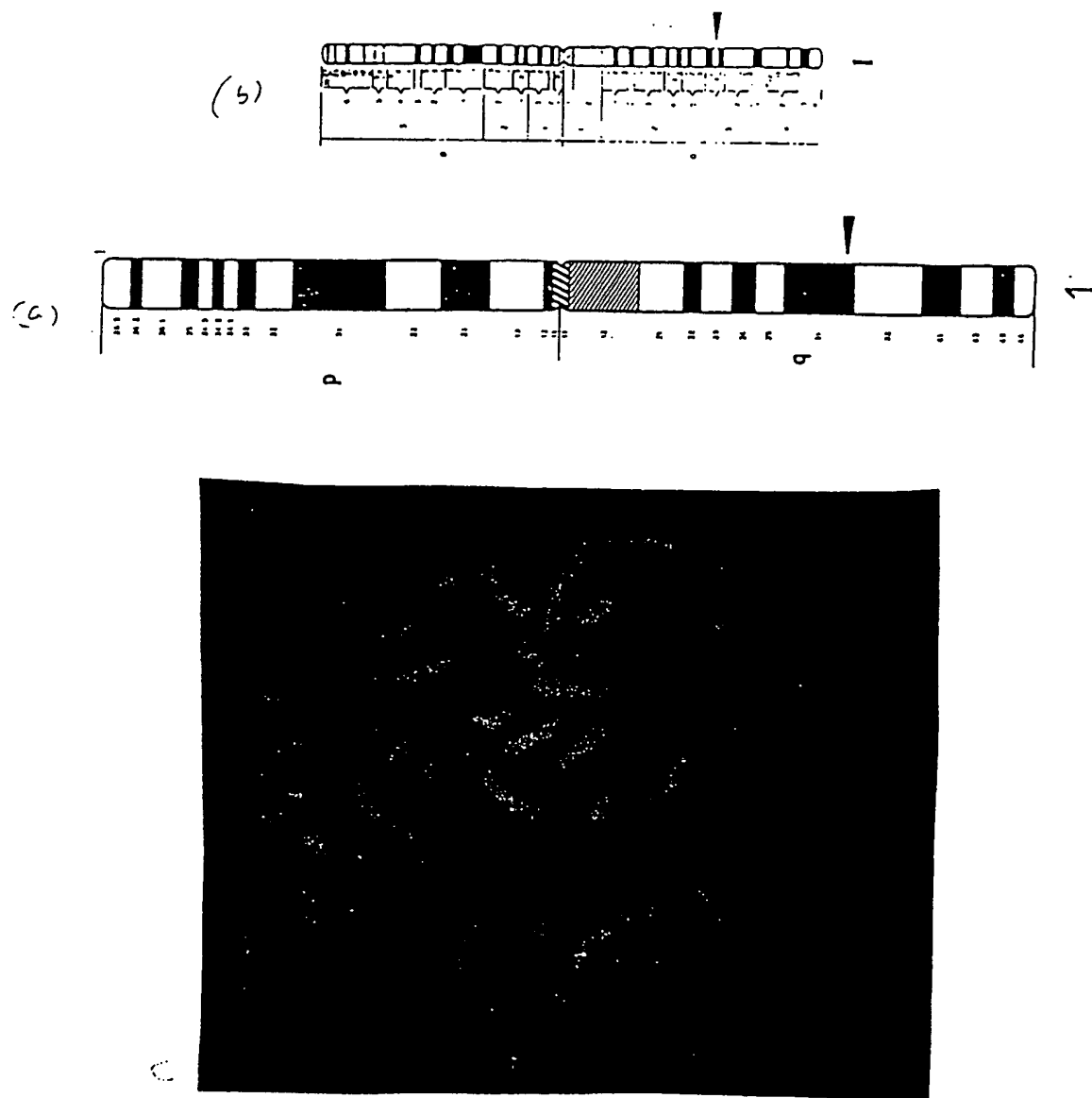


Figure 25a, b, c: Chromosomal localisation of hu-unc-53/l by FISH

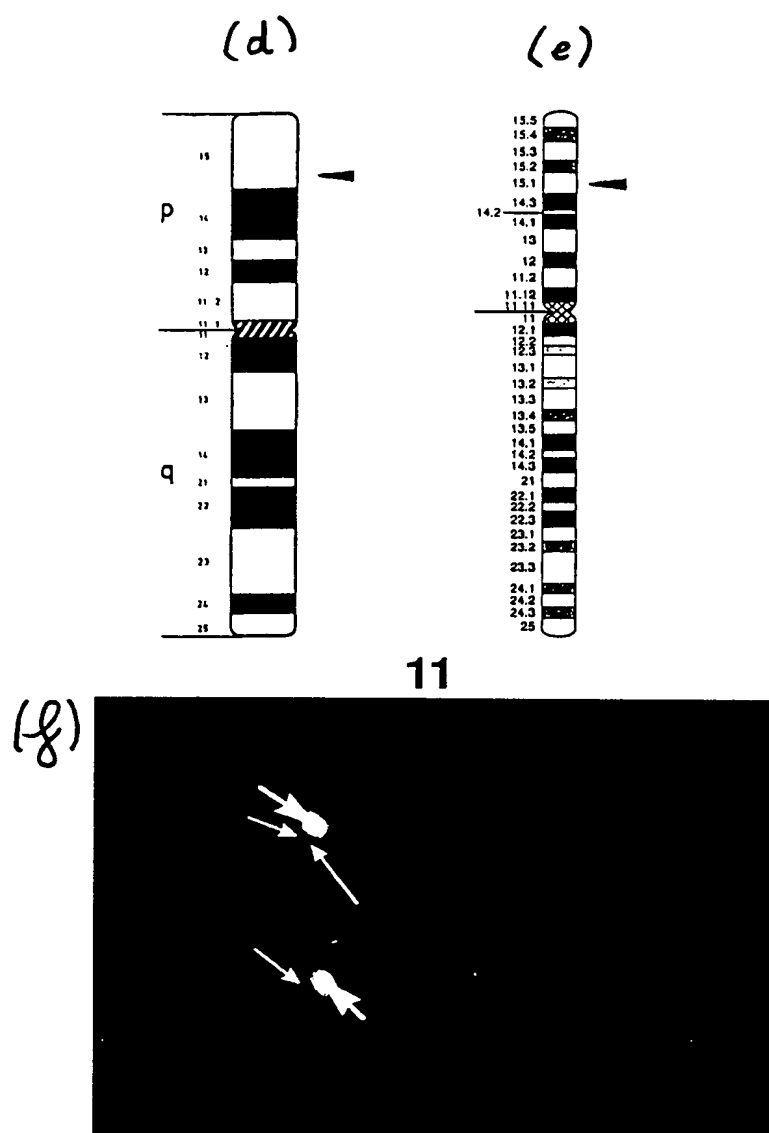


Figure 25d, e, f: Chromosomal localisation of hu-unc-53/2 by FISH

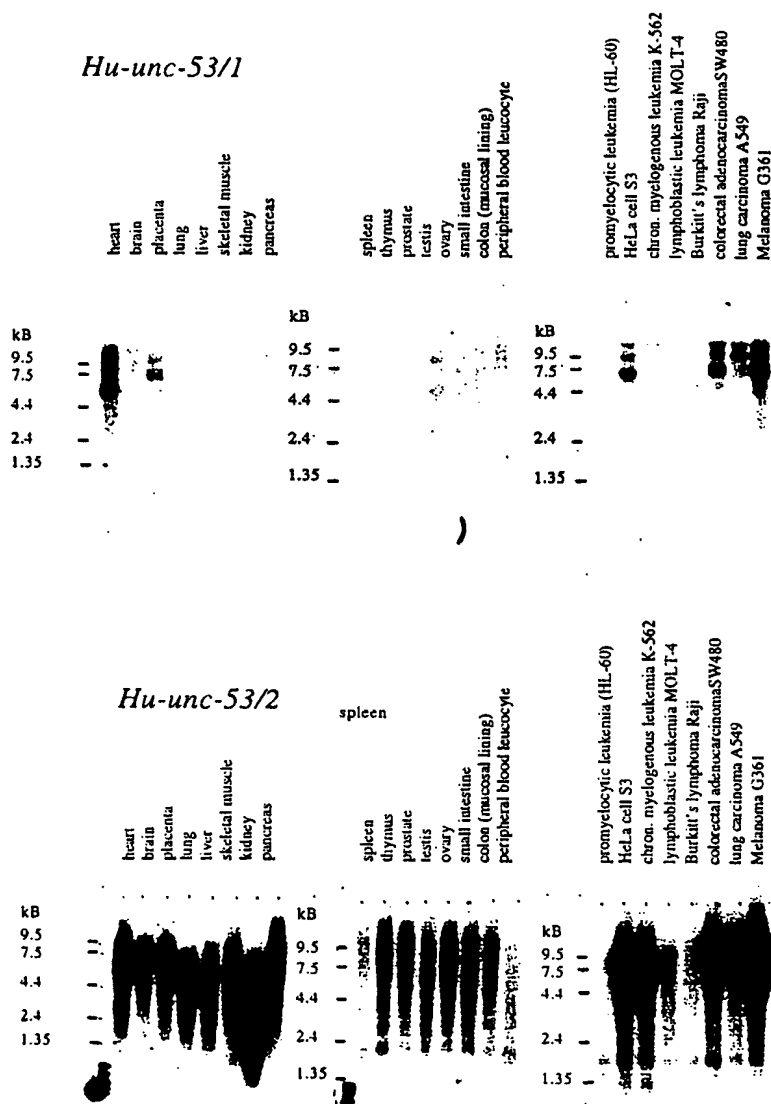


Figure 26 : Expression pattern of *Hu-unc-53/1* and *Hu-unc-53/2* in normal human tissues and cancer cell lines by Northern blotting.

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fig 27 pNP3 Map (1 > 13621) Site and Sequence

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Settings: Linear, Certain Sites Only, Standard Genetic Code

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fig 27 pNP3 Map (1 > 13621) Size and Sequence

Page 2

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fig 27 pNP3 Map (1 > 13621) Site and Sequence

Page 3

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fig 27 pNP3 Map (1 > 13621) Site and Sequence

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FIGURE 28.

SIGNATURE SEQUENCES :

Different signatures can be used to define to identify the UNC-53 gene family :

Aminoacids are listed in one letter code

X equals any aminoacid

X(3,5) equals 3 to 5 X's

(D,E) means D or E at a given position

The signatures should be used to screen a database using a weight matrix of conservative substitutions.

BLOCK A :

GSXLSLASX(3,5)YXSXXEX(4,5)IRXXXR(D,E)LEXXXVXXLTXXXXXXXXLX
XXFEQXL

BLOCK B :

KXKKSXXXXXXXXFXK

BLOCK C :

LARGE FAMILY :

VXXL(K,R)XELX(D,E)(R,K)(D,E)XXLXX(V,I)RL(D,E)XLXXAXXXDXLRE(T,A)X
XXXXXEXXXLKXEXD(R,K)LX

VERTEBRATE FAMILY :

VXXLRXELX(D,E)(R,K)(D,E)MKLTDIRLEALXSAHQLDQLLREXMXNMQXEXX
LKAENDRLK

BLOCK D :

LARGE FAMILY :

W(K,D)X(I,L)DXX(I,V)XX(L,V)F(K,E)XY(I,V,L)XXXDXXXLG(I,L)X(2,3)(D,E)S
(I,V)XGYXI(G,S)(E,H)(L,V,I)(R,K)(R,K)

VERTEBRATE FAMILY :

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S(I,V)XGYSI(G,S)XXKR

BLOCK E :

GXXGXGKS/T

and

F(K,R)MXXXSXX(3,8)GF(I,L,V)(I,L,V)(R,K)Y(I,L,V)(R,K)(R,K)XV(D,E)

and

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L,I)XXXA(R,K)DGX(K,R)XXGXXXX(F,W)EDP

Block F

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fig 29 pEGFPsac (1 > 5100) Site and Sequence

Enzymes : 72 of 146 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

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7p.

Bgl I

TAGTTATTAATAGTAATCAATTACGGGGTCATTAGTTCATAGCCCATATATGGAGTTCGCGTTACATAACTTACGGTAAATGGCCCCCTGGCTGACCG

ATCAATAATTATCATTAGTTAATGCCCCAGTAATCAAGTATCGGGTATATACCTCAAGGCGCAATGTATTGAATGCCATTACCGGGCGGACCGACTGGC

L L I V I N Y G V I S S . P I Y G V P R Y I T Y G K V P A V L T

Aat II

CCCAACGACCCCCGCCCATTGACGTC AATAATGACGTATGTTCCCATAGTAACGCCAATAGGGACTTTCATTGACGTC AATGGGTGGAGTATTTACGGT

GGGTTGCTGGGGGCGGGTAATGTCAGTTATTACTGCATACAAGGGTATCATTCGGGTATCCCTGAAAGGTAAGTGCAGTTACCCACCTCATAAATGCCA

A Q R P P P I D V N N D V C S H S N A N R D F P L T S M G G V F T V

Bgl I Nde I Aat II Bgl I

AAACTGCCCCTTGCGAGTACATCAAGTGTATCATATGCCAAGTACGCCCCCTATTGACGTC AATGACGGTAAATGGCCCCCTGGCATTATGCCAGTA

TTTGACGGGTGAACCGTCATGTAGTTACATAGTATACGGTTCATCGGGGGGATAACTGCAGTTACTGCCATTACCGGGCGGACCGTAATACGGGTCAT

N C P L G S T S S V S Y A K Y A P Y . R Q . R . M A R L A L C P V

SnaB I Nco I

CATGACCTTATGGGACTTTCCTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGATGCGGTTTGGCAGTACATCAATGGGCGTGGA

GTA CTGGAATACCTGAAAGGATGAACCGTCATGTAGATGCATAATCAGTAGCGATAATGGTACCAC TAGCCAAAACCGTCATGTAGTTACCGGCACCT

H D L M G L S Y L A V H L R I S H R Y Y H G D A V L A V H Q V A V

Aat II

TAGCGGTTTGACTCAGCGGGATTTC AAGTCTCCACCCCATTGACGTC AATGGGAGTTTGTTTGGCACCAAAATCAACGGGACTTTC AAAATGTCGTA

ATCGCCAAACTGAGTGCCCTAAAGGTT CAGAGGTGGGGTAAGTGCAGTTACCTCAAACAAAACCGTGGTTT TAGTTGCCCTGAAAGGTTTACAGCAT

I A V . L T G I S K S P P H . R Q V E F V L A P K S T G L S K M S .

Nhe I Eca7

ACAAC TCCGCCCCATTGACGCA AATGGGCGGTAGGCGTG TACGGTGGGAGGTC TATATAAGCAGAGCTGGTTTAGTGAACCGTCAGATCCGCTAGCGCTA

TGTTGAGGGCGGGTAAC TGCGTTTACCCGCCATCCGCACATGCCACCCTCCAGATATATTCGCTCGACCAAATCACTTGGCAGTCTAGGCGATCGCGAT

O L R P I D A N G R . A C T V G G L Y K O S V F S E P S D P L A L

Nco I

CCGGTGGCCACCATGGTGAGCAAGGCGGAGGAGCTGTTTACCGGGGTGGTGCCCATCTGGTTCGAGCTGGACGGGACGTAACGGGCCACAAGTTCAGCG

GGCCAGCGGTGGTACCACCTCGTTCCGGCTCCTCGACAAGTGGCCCCACACGGGTAGGACCAGCTCGACCTGCCCGTGCATTTCGCCGTGTTCAAGTCGG

P V A T M V S K G E E L F T G V V P I L V E L D G D V N G H K F S

TGTCGGGCGAGGGCGAGGGCGATGCCACCTACGGCAAGCTGACCTGAAGTTCTATCGACCAACCGGCAAGCTGCCCGTGCCCTGGGCCACCTCGTGAC

ACAGGCGGCTCCCGCTCCCGCTACGGTGGATGCCGTTTCGACTGGGACTTCAAGTAGACGTGGTGGCCGTTCGACGGGACGGGACCGGGTGGGAGCACTG

V S G E G E G D A T Y G K L T L K F I C T T G K L P V P V P T L V T

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fig 29 pEGFPsac (1 > 5100) Site and Sequence

Page 2

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GTGGGACTGGATGCCGACGTCACGAAGTCGGCGATGGGGCTGGTGTACTTCGTCTGCTGAAGAAGTTCAGGCGGTACGGGCTTCCGATGCAGGTCTCTC 900

.....
KspI
.....
T L T Y G V Q C F S R Y P D H M K Q H D F F K S A M P E G Y V Q E

CGCACCATCTTCTTCAAGGACGACGGCAACTACAAGACCCGCGCGAGGTGAAGTTCGAGGGCGACACCCTGGTGAACCGCATCGAGCTGAAGGGCATCG 1000
GCGTGGTAGAAGAAGTTCCTGCTGCCGTGTATGTTCTGGGCGCGGCTCCACTTCAAGCTCCCGCTGTGGGACCAC TTGGCGTAGCTCGACTTCCCGTAGC

.....
R T I F F K D D G N Y K T R A E V K F E G D T L V N R I E L K G I

ACTTCAAGGAGGACGGCAACATCTGGGGCACAAGCTGGAGTACAACACAGCCACAACGTCTATATCATGGCCGACAAGCAGAAGAACGGCATCAA 1100
TGAAGTTCTCTGCGGTTGTAGGACCCGTTGTCGACCTCATGTTGATGTTGTCGGTGTTCAGATATAGTACCGGCTGTTCTGCTCTTCTGCGGTAGTT

.....
D F K E D G N I L G H K L E Y N Y N S H N V Y I M A D K Q K N G I

GGTGAAC TTCAAGATCCGCCACAACATCGAGGACGGCAGCGTGCAGCTCGCCGACCCTACCAGCAGAACACCCCATCGGCGACGGCCCCGTGCTGCTG 1200
CCACTTGAAGTTCTAGGCGGTGTGTAGCTCCTGCCGTCGCACGTCGAGCGGCTGGTGATGGTCGCTTGTGGGGGTAGCCGCTGCCGGGGCACGACGAC

.....
V N F K I R H N I E D G S V Q L A D H Y Q Q N T P I G D G P V L L

CCCGACAACCACTACCTGAGCACCCAGTCCGCCCTGAGCAAGACCCCAACGAGAAGCGGATCACATGGTCC TGCTGGAGTTCTGTGACCSCCGCCGGGA 1300
GGGCTGTTGGTGATGGACTCGTGGGTCAGGCGGGACTCGTTTCTGGGGTTGCTCTTCGCGCTAGTGTTACAGGACGACCTCAAGCAC TGGCGGCGGCCCT

.....
P D N H Y L S T Q S A L S K D P N E K R D H M V L L E F V T A A G

.....
BspM II Bgl II Asu II EcoN I
TCACTCTCGGATGGACGAGCTGTACAAGTCCGGACTCAGATCTACGTCAAATGTAGAATTGATACCAATCTACACGGATTGGGCCAATCGGCACCTTTC 1400
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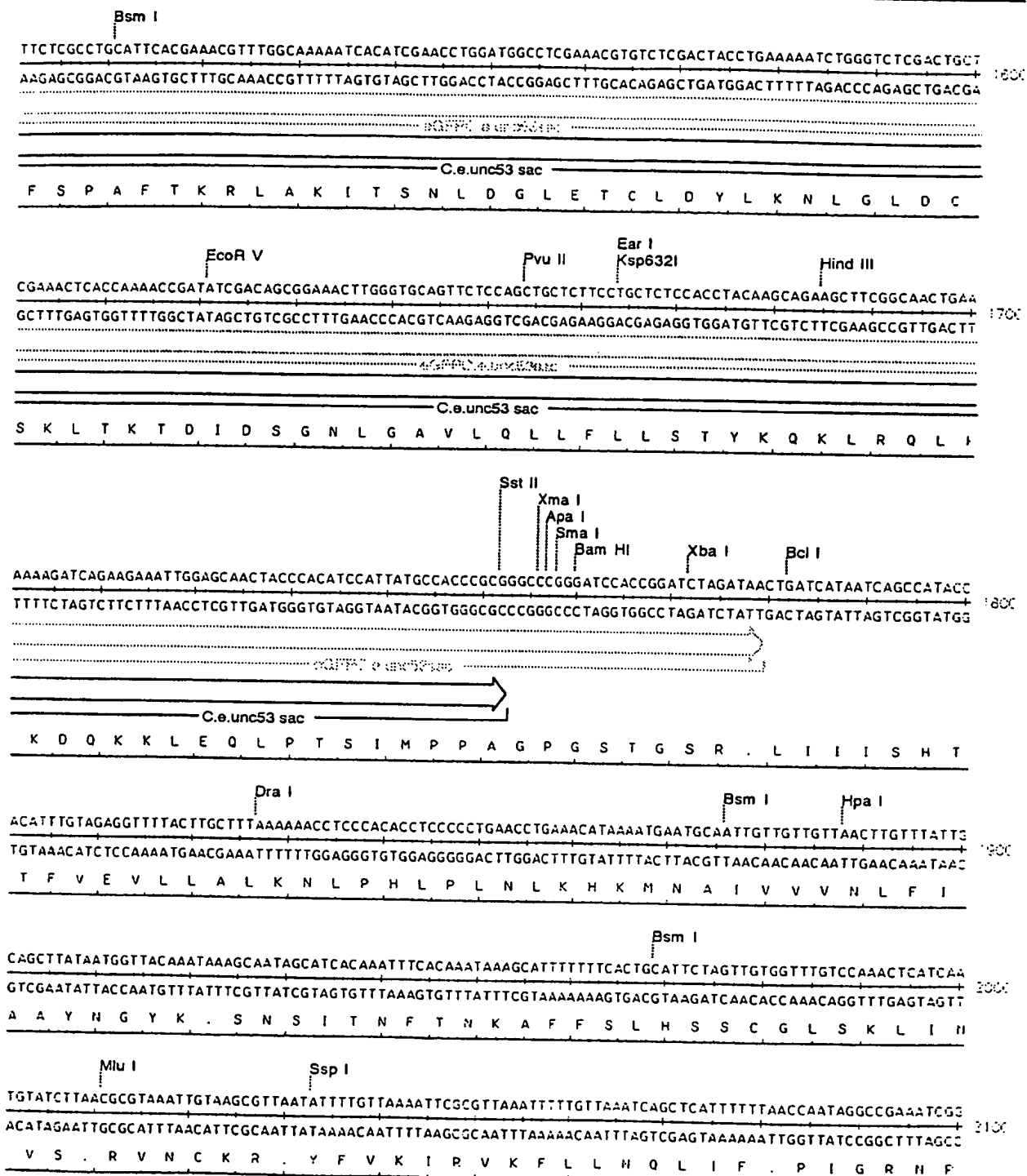
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C.e.unc53 sac
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.....
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CTTCCCGTCGAATAGTTTTCAGCTAATCCCTATAAAGGTTACTAAAAGCGCTATAGTGCACCAAGAGTCGAATAATTACACTAGCAAGGCTAGTTGCTT

.....
C.e.unc53 sac
K G S L S K S I R D I S N D F R D Y R L V S Q L I N V I V P I N E

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lig 29 pEGFPsac (1 > 5100) Site and Sequence

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Bsr I

CAAAATCCCTTATAAATCAAAGAATAGACCGAGATAGGGTTGAGTGTGTTCAGTTTGGAAACAGAGTCCACTATTAAGAAGCTGGACTCCAACGTC
GTTTTAGGGAATATTTAGTTTTCTTATCTGGCTCTATCCCACTCACAACAAGGTCAAACCTGTTCTCAGGTGATAATTTCTTGACCTGAGGTTGCAG 3200
Q N P L . I K R I D R D R V E C C S S L E Q E S T I K E R G L Q R

Dra III

AAAGGGCGAAAAACCGTCTATCAGGGCGATGGCCCACTACGTGAACCATCACCTAATCAAGTTTTTGGGGTCGAGGTGCCGTAAAGCACTAAATCGGA
TTTCCCGCTTTTGGCAGATAGTCCCGCTACCGGGTGATGCACCTGGTAGTGGGATTAGTTCAAAAAACCCAGCTCCACGGCATTTCTGTGATTTAGCCT 3300
Q R A K N R L S G R V P T T . T I T L I K F F G V E V P . S T K S E

Nae I

ACCCTAAAGGAGCCCCGATTTAGAGCTTGACGGGGAAGCCGGCGAACGTGGCGAGAAAGGAAGGAAGAAAGCGAAAGGAGCGGGCGCTAGGGCGCT
TGGGATTTCCCTCGGGGGCTAAATCTCGAACTGCCCTTTCGGCCGCTTGACCCGCTCTTTCCCTTCCTTTCTCGCTTTCCTCGCCCGCATCCCGCGA 3400
P . R E P P I . S L T G K A G E R G E K G R E E S E R S G R . G A

Ksp I

GGCAAGTGTAGCGGTACGCTGCGCGTAACCACCACACCCCGCCGCTTAATGCGCGCTACAGGGCGCGTCAGGTGGCACTTTTCGGGAAATGTGGC
CCGTTACATCGCCAGTGGACGCGCATTTGGTGGTGTGGGCGGCGGAATTACGCGCGATGTCCCGCGCAGTCCACCGTGAAAAGCCCTTTACACGGC 3500
G K C S G H A A R N H H T R R A . C A A T G R V R V H F S G K C A

BspH I

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CCTTGGGGATAAACAAATAAAAGATTTATGTAAGTTTATACATAGGCGAGTACTCTGTTATTGGGACTATTACGAAGTTATTATAACTTTTCTTCT 3600
R N P Y L F I F L N T F K Y V S A H E T I T L I N A S I I L K K E E

Ssp I

Ear I

Ksp632I

OxaN I

Pvu II

Sph I

Ava III

Nsi I

GTCTGAGGCGGAAGAACCAGCTGTGAATGTGTGTCAGTTAGGGTGTGGAAGTCCCCAGGCTCCCCAGCAGGCAGAAGTATGCAAGCATGCATCTC
CAGGACTCCGCCTTCTTGGTCGACACCTTACACACAGTCAATCCACACCTTTCAGGGTCCGAGGGTCTGTCCTTTCATACGTTTCGTACGTAGAG 3700
S . G G K N Q L V N V C Q L G C G K S P G S P A G R S M Q S M H L

Sph I

Ava III

Nsi I

AATTAGTCAGCAACCAGGTGTGGAAGTCCCCAGGCTCCCCAGCAGGCAGAAGTATGCAAGCATGCATCTCAATTAGTCAGCAACCATAGTCCCGGCC
TTAATCAGTCGTTGGTCCACACCTTTCAGGGTCCGAGGGTCTGTCCTTTCATACGTTTCTGACGTAGAGTTAATCAGTCGTTGGTATCAGGCGCGG 3800
N . S A T R C G K S P G S P A G R S M Q S M H L N . S A T I V P P

Bsr I

Nco I

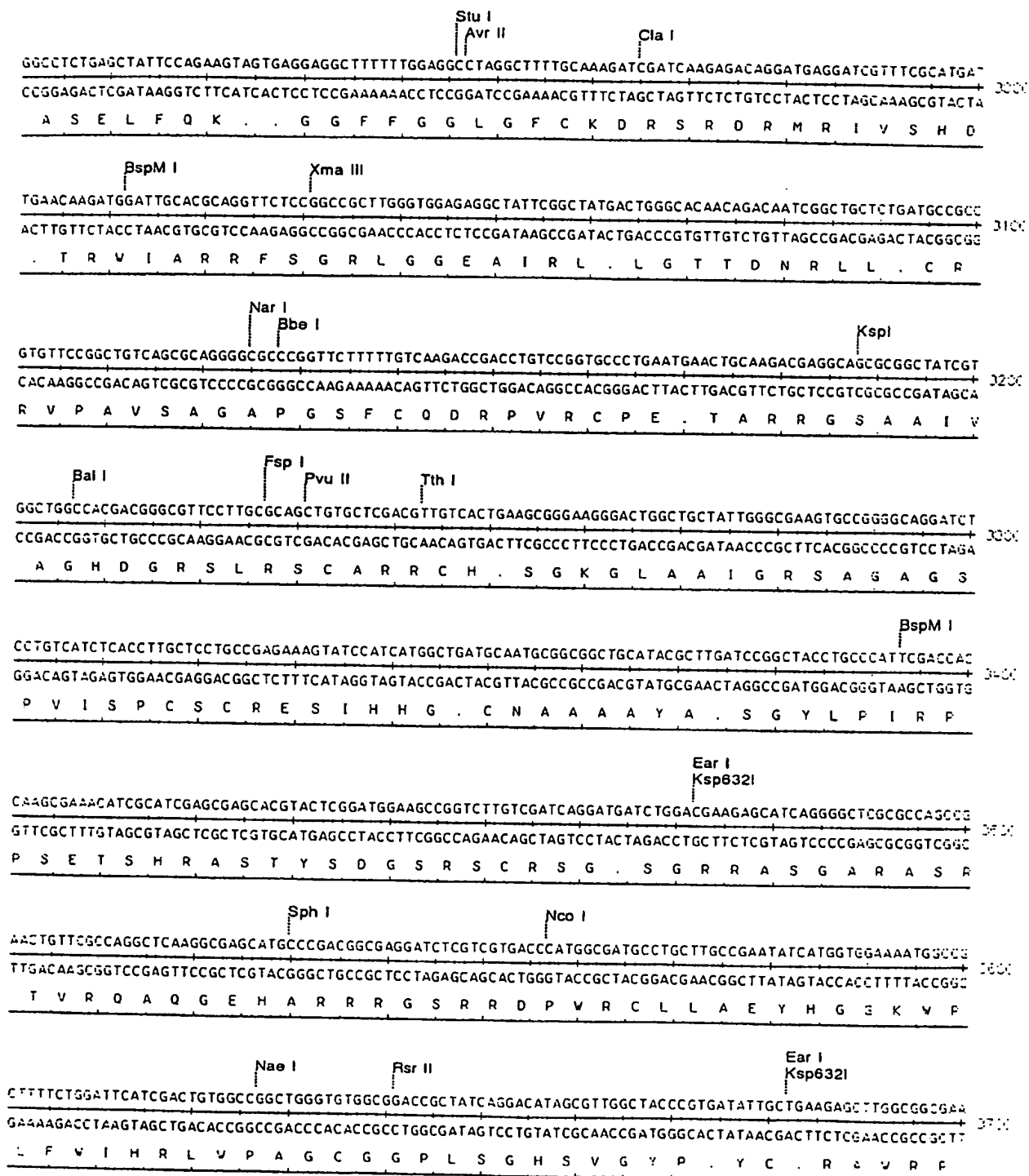
Bgl I

Sfi I

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ATTGAGGCGGGTAGGCGGGGATTGAGGCGGGTCAAGGCGGGTAAGAGGCGGGTACCGACTGATTAATAAATAAATACGTCCTCGGCTCCGCGCGGAG 3900
L T P P I P P L T P P S S A H S P P H G . L I F F I Y A E A E A A

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fig 29 pEGFPsac (1 > 5100) Site and Sequence

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fig 29 pEGFPsac (1 > 5100) Site and Sequence

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TGGGCTGACCGCTTCCTCGTGCTTTACGGTATCGCCGCTCCCGATTGCGAGCGCATCGCCTTCTATCGCCTTCTTGACGAGTTCTTCTGAGCGGGACTCT
ACCCGACTGGCGAAGGAGCACGAAATGCCATAGCGGCGAGGGCTAAGCGTCCGCTAGCGGAAGATAGCGGAAGAAGTCTCAAGAAGACTCGCCCTGAGA 3800
M G . P L P R A L R Y R R S R F A A H R L L S P S . R V L L S G T L

Asu II BspM I
GGGGTTCGAAATGACCGACCAAGCGACGCCAACCTGCCATCAGAGATTTCGATTCCACCGCCGCCCTTCTATGAAAGGTTGGGCTTCGGAATCGTTTTTC
CCCCAAGCTTTACTGGCTGGTTCGCTGCGGGTTGGACGGTAGTGCTCTAAAGCTAAGGTGGCGGCGGAAGATACTTTCCAACCCGAAGCCTTAGCAAAAG 3900
G F E M T D Q A T P N L P S R D F D S T A A F Y E R L G F G I V F

Nae I KspI Avr II
CGGGACGCCGCTGGATGATCTCCAGCGCGGGGATCTCATGCTGGAGTTCTTCGCCACCCCTAGGGGGAGGCTAACTGAAACACGGAAGGAGACAATAC
GCCCTGCGGCGGACCTACTAGGAGGTGCGGCCCTTAGAGTACGACCTCAAGAAGCGGGTGGGATCCCGCTCCGATTGACTTTGTGCTTCTCTGTATTG 4000
R D A G W M I L O R G D L M L E F F A H P R G R L T E T R K E T I

KspI
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GCCCTCCTTGGGCGGATACTGCCGTTATTTTCTGTCTTATTTGCGTGCCACAACCCAGCAACAAGTATTTGCGCCCAAGCCAGGGTCCCGACCGT 4100
P E G T R A M T A I K R O N K T H G V G S F V H K R G V R S Q G W H

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S V D T P P R P H W G Q Y A R V S S F S P P H P P S S G E G P G L

AlwI OxaI PraI PraI
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CGTCGGTTCAGCCCCGCGCTCCGGGACGGTATCGGAGTCCAATGAGTATATATGAAATCTAACTAAATTTGAAGTAAAAATTAATTTTCTAGATCC 4300
A A N V G A A G P A I A S G Y S Y I L . I D L K L H F . F K R I .

BspH I
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V K I L F D N L M T K I P . R E F S F H . A S D P V E K I K G S S .

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D P F F L R V I C C L Q T K K P P L P A V V C L P D Q E L P T L F

BsrI
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GGCTTCCATTGACCGAAGTCGCTCGCGCTATGGTTTATGACAGGAAGATCACATCGGCATCAATCCGGTGGTGAAGTTCTTGAGACATCGTGCGGAT 4600
P K V T G F S R A Q I P N T V L L V . P . L G H H F K N S V A P P

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fig 29 pEGFPsac (1 > 5100) Site and Sequence

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AlwI I
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GTATGGAGCGAGACGATTAGGACAATGGTCACCGACGACGGTCACCGCTATTCAGCACAGAATGGCCCAACCTGAGTTCTGCTATCAATGSCCTATTCCG
T Y L A L L I L L P V A A A S G D K S C L T G L D S R R . L P D K A 4700

ApaI I
GCAGCGGTCGGGCTGAACGGGGGTTCTGTGCACACAGCCAGCTTGGAGCGAACGACCTACACCGAACTGAGATACCTACAGCGTGAGCTATGAGAAAGC
CGTCGCCAGCCGACTTGCCCCCAAGCACGTGTGTCGGGTCGAACCTCGCTTGGATGTGGCTTGACTCTATGGATGTCGCACTCGATACTCTTTCC
Q R S G . T G G S C T Q P S L E R T T Y T E L R Y L Q R E L . E S 4800

GCCACGCTTCCC GAAGGGAGAAAGCGGACAGGTATCCGTAAGCGGCAGGGTCGGAACAGGAGAGCGCACGAGGGAGCTTCCAGGGGGAAACGCTGGT
CGGTGCGAAGGGCTTCCCTCTTTCCGCTGTCCATAGGCCATTGCGGTCGCCAGCCTTGCTCTCGCGTGCTCCCTCGAAGGTCCCCCTTTGCGGACCA
A T L P E G R K A D R Y P V S G R V G T G E R T R E L P G G N A W 4900

ATCTTTATAGTCTGTGCGGGTTTCGCCACCTCTGACTTGAGCGTCGATTTTGTGATGCTCGTCAGGGGGCGGAGCCTATGGAAAAACGCCAGCAACGC
TAGAAATATCAGGACAGCCAAAGCGGTGGAGACTGAATCGCAGCTAAAAACACTACGAGCAGTCCCCCGCCTCGGATACCTTTTTCGGGTCGTGCG
Y L Y S P V G F R H L . L E R R F L . C S S G G R S L V K N A S N A 5000

Ava III
Nsi I
GGCCTTTTACGGTTCTTGCCCTTTTGTGCGCCTTTTGTTCACATGTTCTTTCTGCGTTATCCCTGATTCTGTGGATAACCGTATTACCGCCATGCAT
CCGGAATAATGCCAAGGACCGGAAACGACCGGAAACGAGTGTAAGAAGAGGACGCAATAGGGGACTAAGACACCTATTGGCATAATGGCGGTACGTA
A F L R F L A F C V P F A H M F F P A L S P D S V D N R I T A M H 5100

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fig 30 pEGFP72 (1 > 9697) Site and Sequence

Enzymes : 72 of 146 enzymes (Filtered)

Settings: Linear, Certain Sites Only, Standard Genetic Code

Page 1

Page 8

16p

Bgl I

100

L L I V I N Y G V I S S . P I Y G V P R Y I T Y G K V P A V L T

Aat II

200

A Q R P P P I D V N N D V C S H S N A N R D F P L T S M G G V F T V

Bgl I

Nde I

Aat II

Bgl I

300

N C P L G S T S S V S Y A K Y A P Y . R Q . R . M A R L A L C P V

SnaB I

Nco I

400

H D L M G L S Y L A V H L R I S H R Y Y H G D A V L A V H Q V A V

Aat II

500

I A V . L T G I S K S P P H . R Q V E F V L A P K S T G L S K M S .

Nhe I

Eco7

600

Q L R P I D A N G R . A C T V G G L Y K Q S V F S E P S D P L A L

Nco I

700

P V A T M V S K G E E L F T G V V P I L V E L D G D V N G H K F S

eGFP.C.e.unc53

800

V S G E G E G D A T Y G K L T L K F I C T T G K L P V P W P I L V T

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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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900

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KspI
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1100

eGFP.C.e.unc53
D F K E D G N I L G H K L E Y N Y N S H N V Y I M A D K Q K N G I A

GGTGAAC TTCAAGATCCGCCACAACATCGAGGACGGCAGCGTGCAGCTCGCCGACCACTACCAGCAGAACACCCCATCGGCGACGGCCCGTGTCTGTG
CCACTTGAAGTTC TAGGCGGTGTTGTAGCTCCTGCCGTCGCAGCTCGAGCGGCTGGTGATGGTCTCTTGTGGGGTAGCCGCTGCCGGGGCACGACGAC
1200

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V N F K I R H N I E D G S V Q L A D H Y Q Q N T P I G D G P V L L

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eGFP.C.e.unc53
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BspM II Bgl II
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1400

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C.e.unc53
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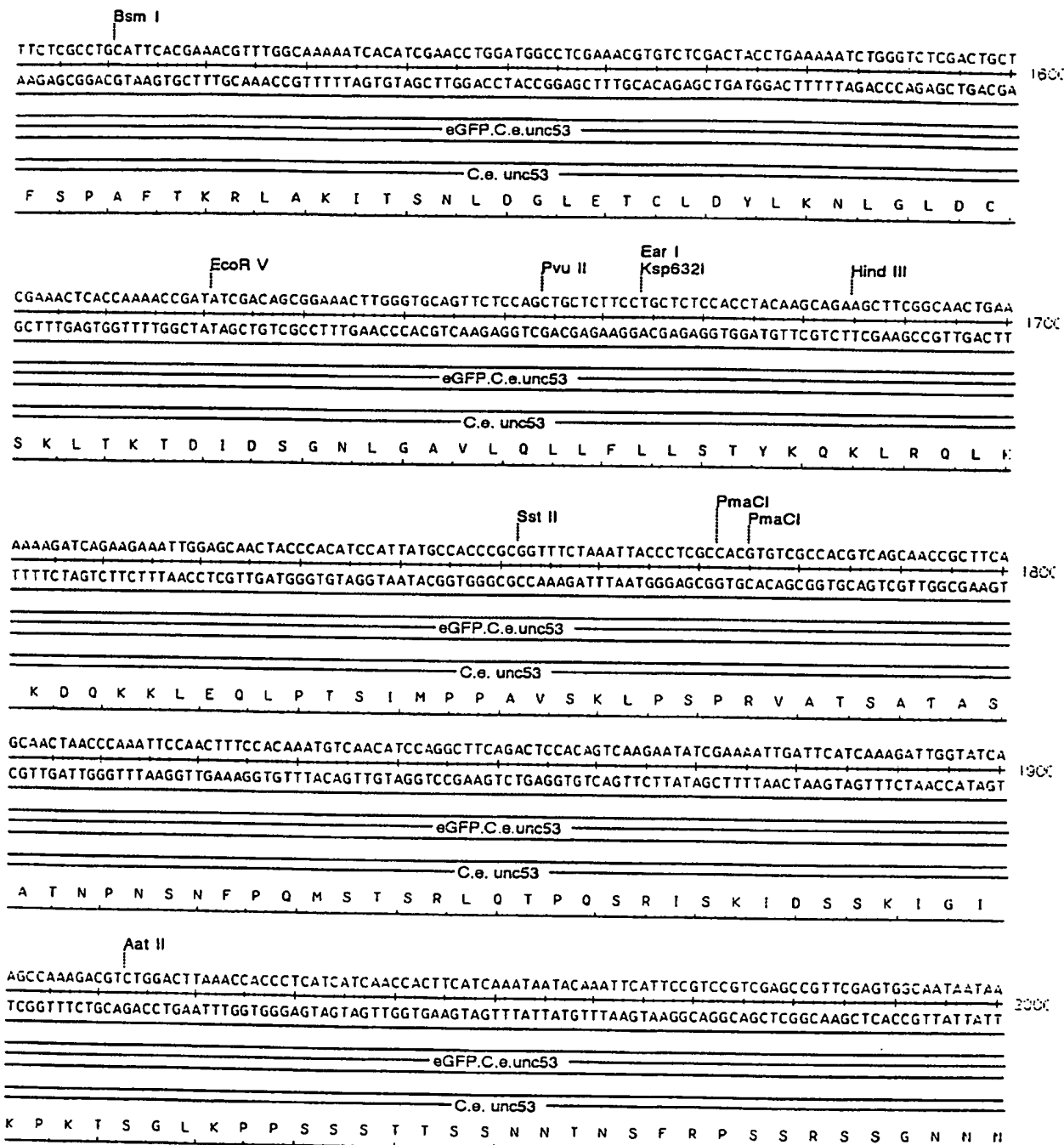
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1500

eGFP.C.e.unc53

C.e.unc53
K G S L S K S I R D I S H D F R D Y R L V S Q L I N V I V P I N E

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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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Ksp632I

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ACAACCGAGCTGCTATAGGTGTAGACGCTTCTCGAATCTTAGTAGTAGTTGCATGTCGAGATAAAGCTTAGATTTGGCTGGATGGAGGGTTGAGGTTTTT

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-----C.e. unc53-----
V G S T I S T S A K S L E S S S T Y S S I S N L N R P T S Q L Q K

Xba I Nhe I

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GGAAGATCTGGTGTGTTGGGTCGATCAAGCACAACGATGTTGATGTTTTAGCCTTCGAGTTTCGATCGGCGAGGCTTTCGGCACTCGTGGGGTTTTGAAC

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-----C.e. unc53-----
P S R P Q T Q L V R V A T T T K I G S S K L A A P K A V S T P K L

Bsm I

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A S V K T I G A K Q E P D N S G G G G G G M L K L K L F S S K N P S

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AAGGAGTAGCTTATCGGGTGTGGATGCTCTTTCCGCGCGCCACGAGTTGTTGTTTGAACAGCTTTTAGCGACGGGGTCACTTTTCACGGGACTTC

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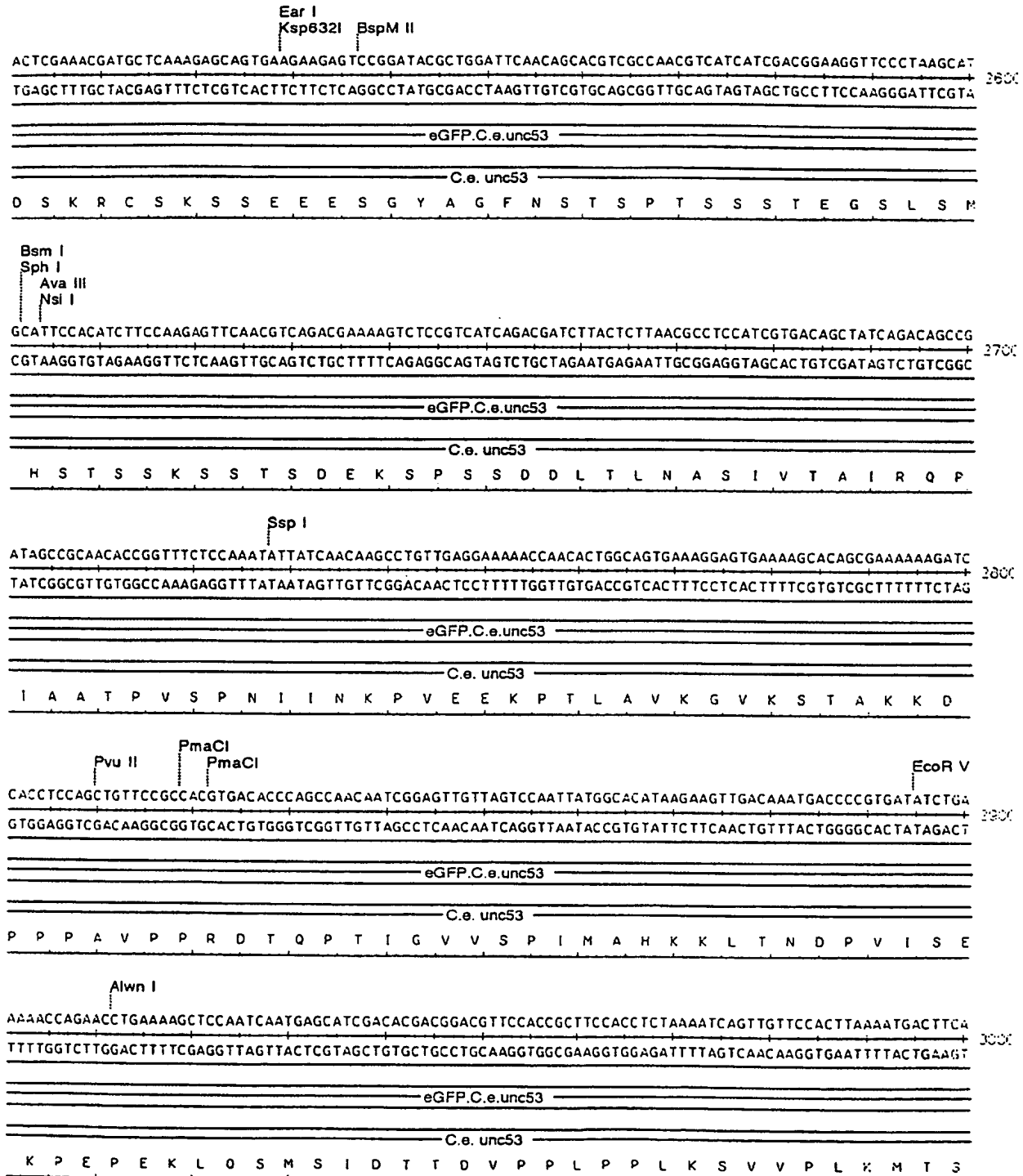
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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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fig 30 pEGFP72 (1 > 9697) Site and Sequence

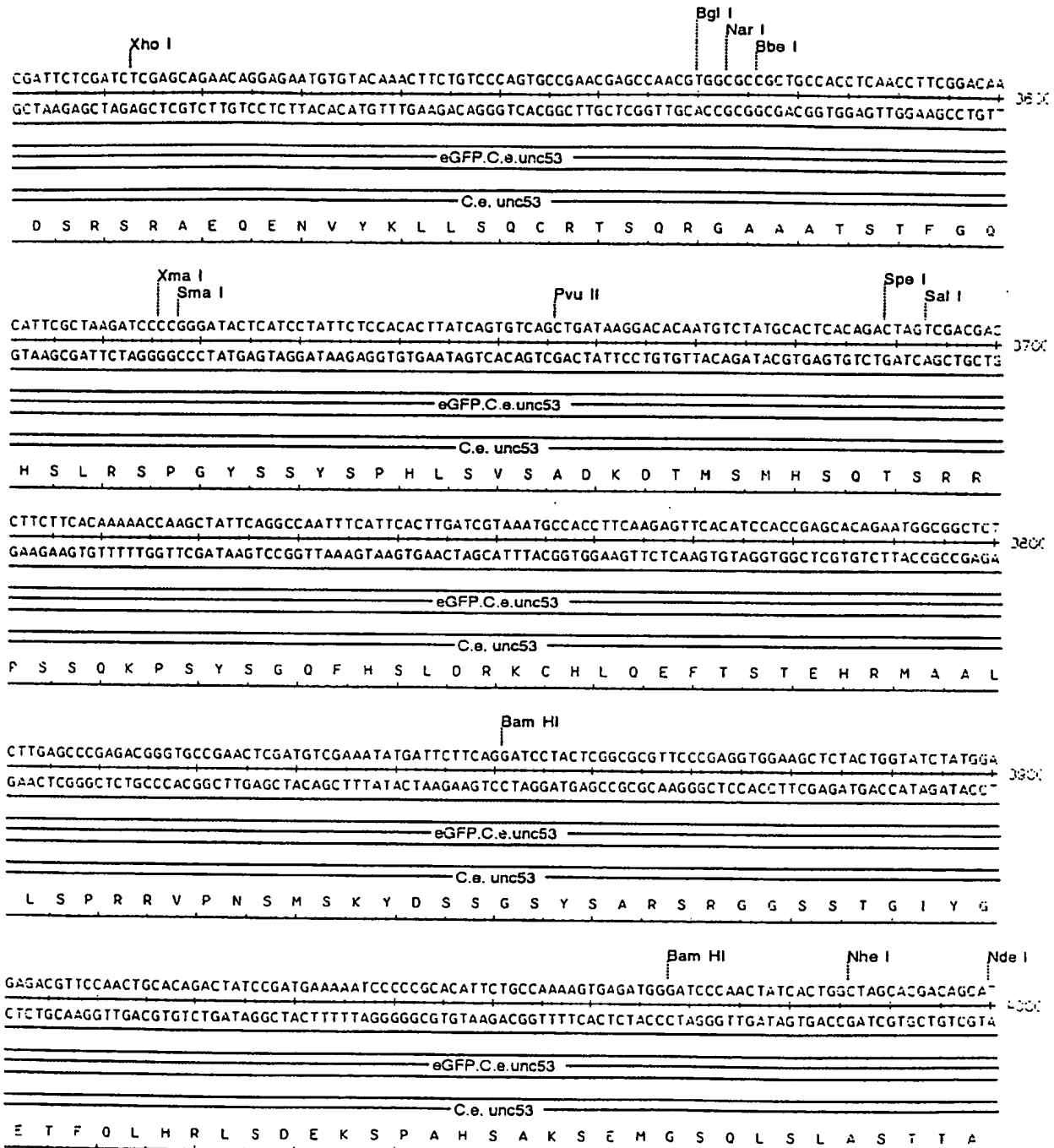
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Spl
Spl

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-----C.e. unc53-----
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-----C.e. unc53-----
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-----C.e. unc53-----
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Asu II Sst I BspM II
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-----eGFP.C.e.unc53-----
-----C.e. unc53-----
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-----eGFP.C.e.unc53-----
-----C.e. unc53-----
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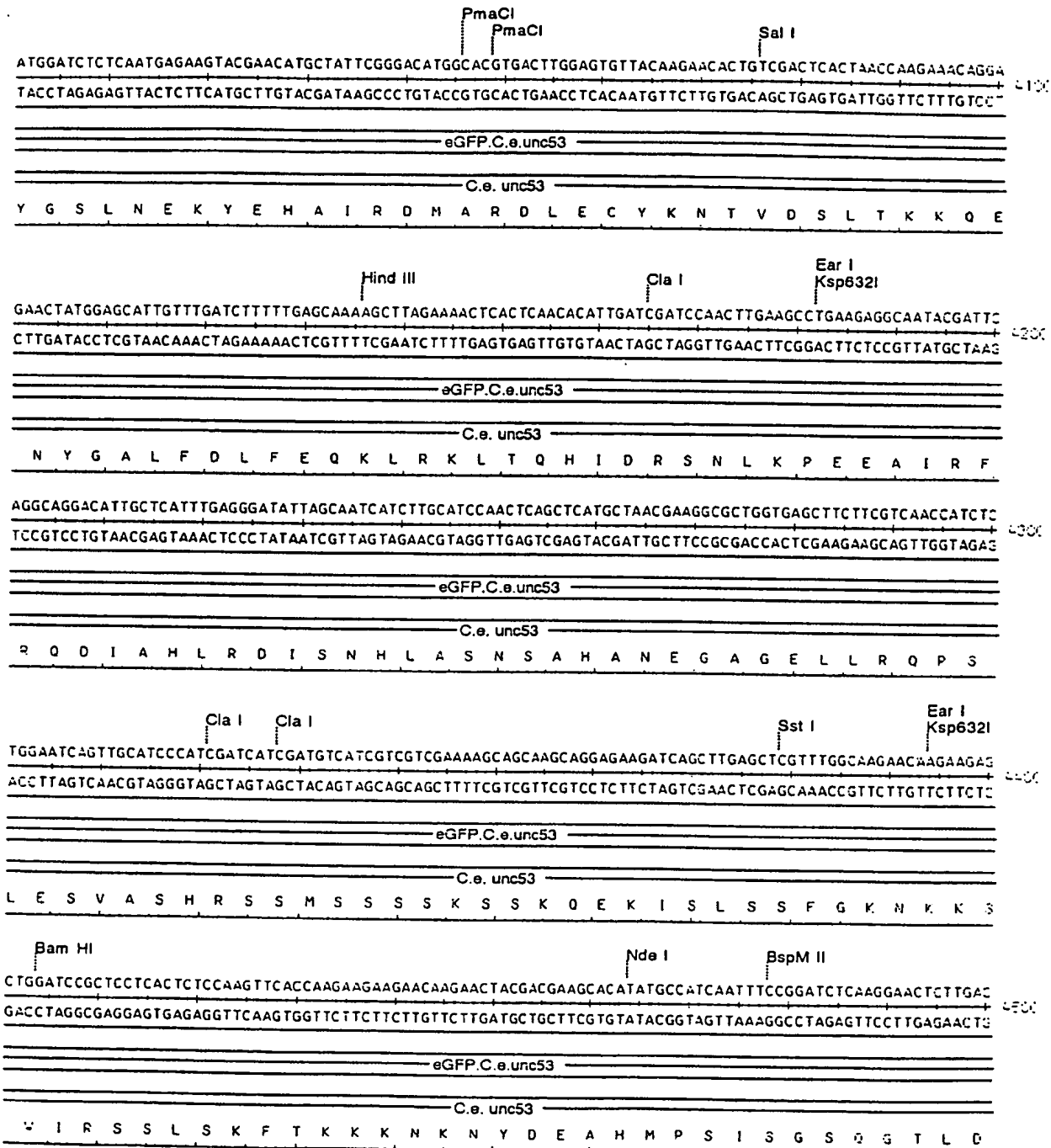
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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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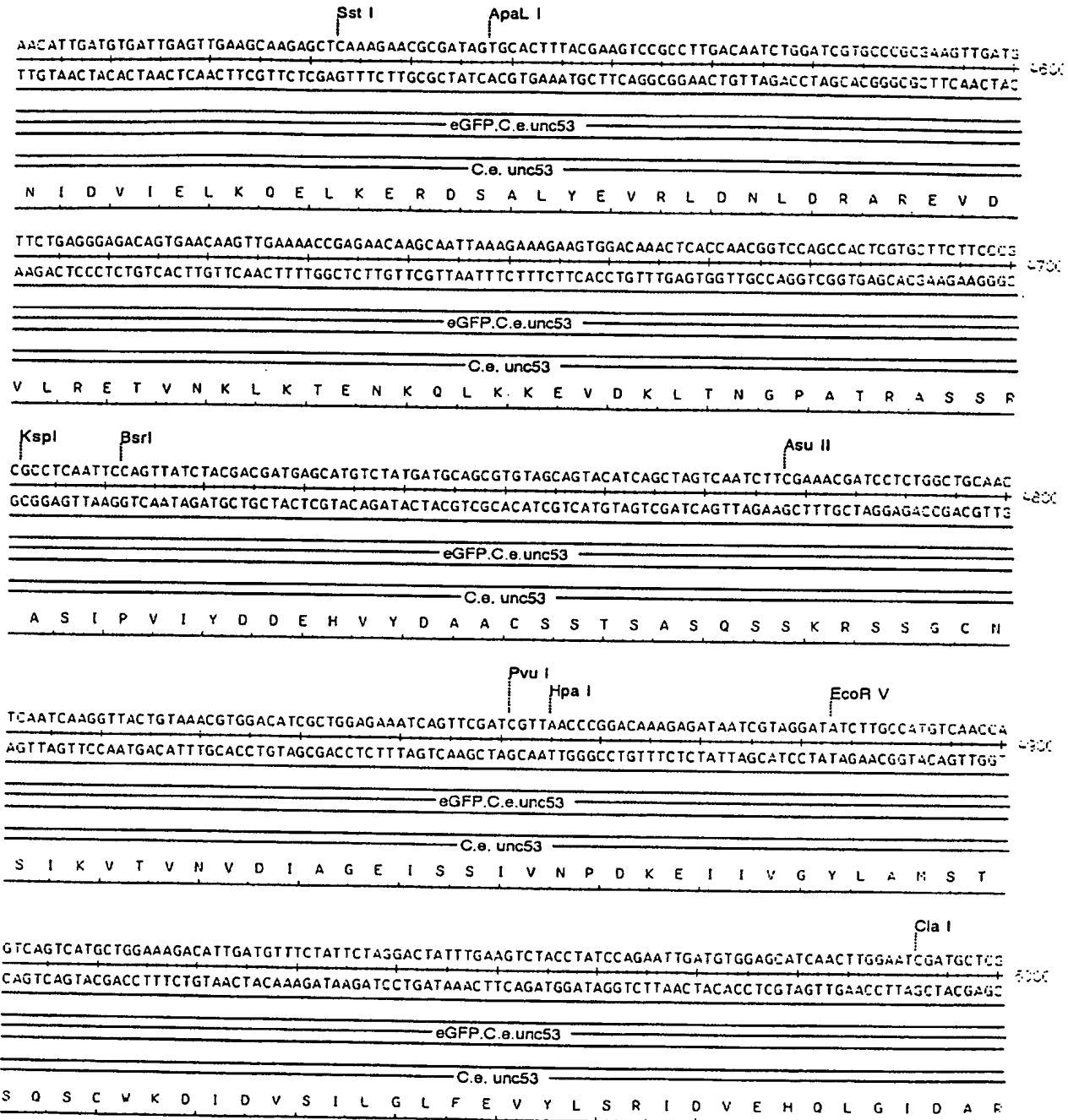
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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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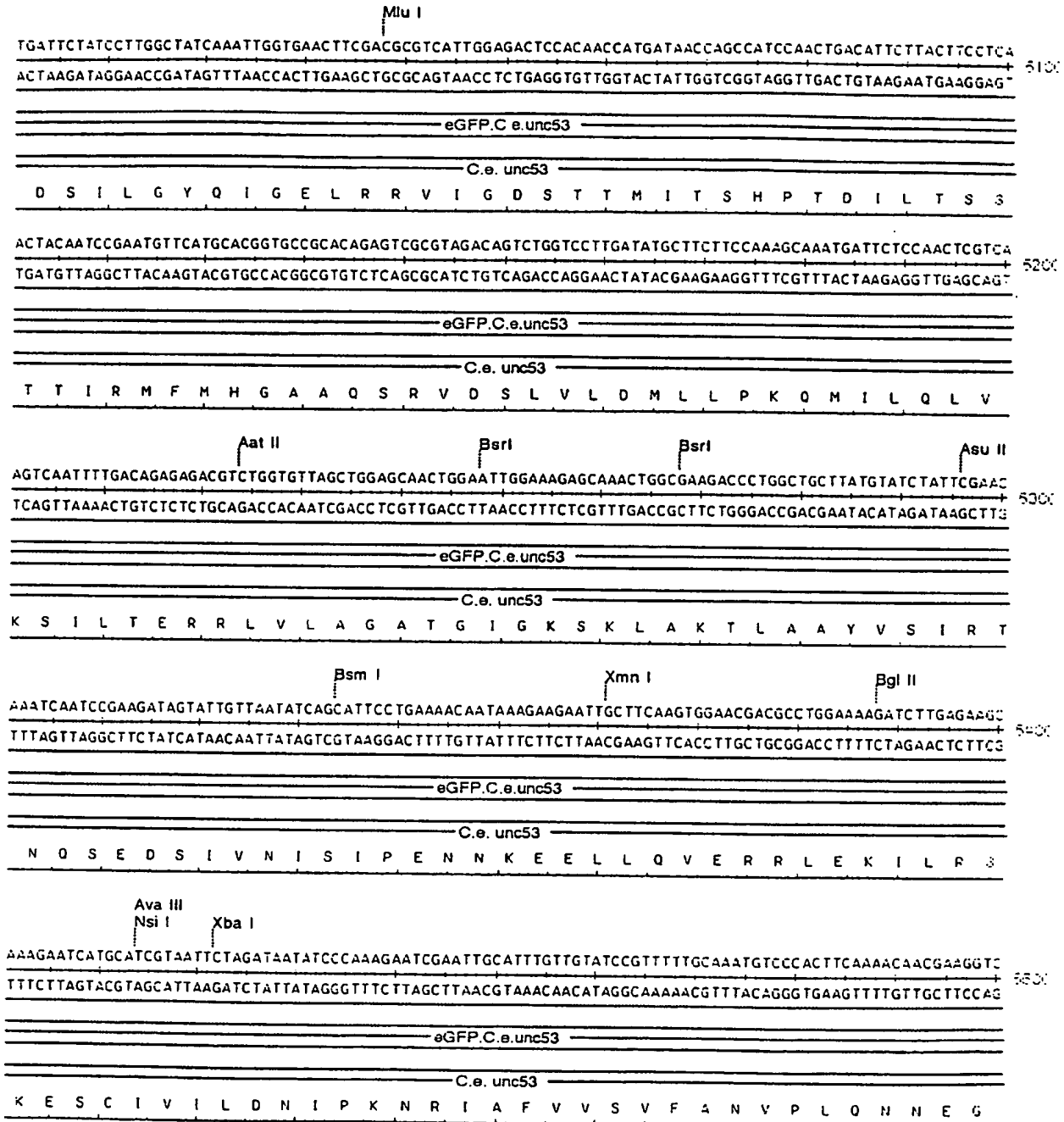
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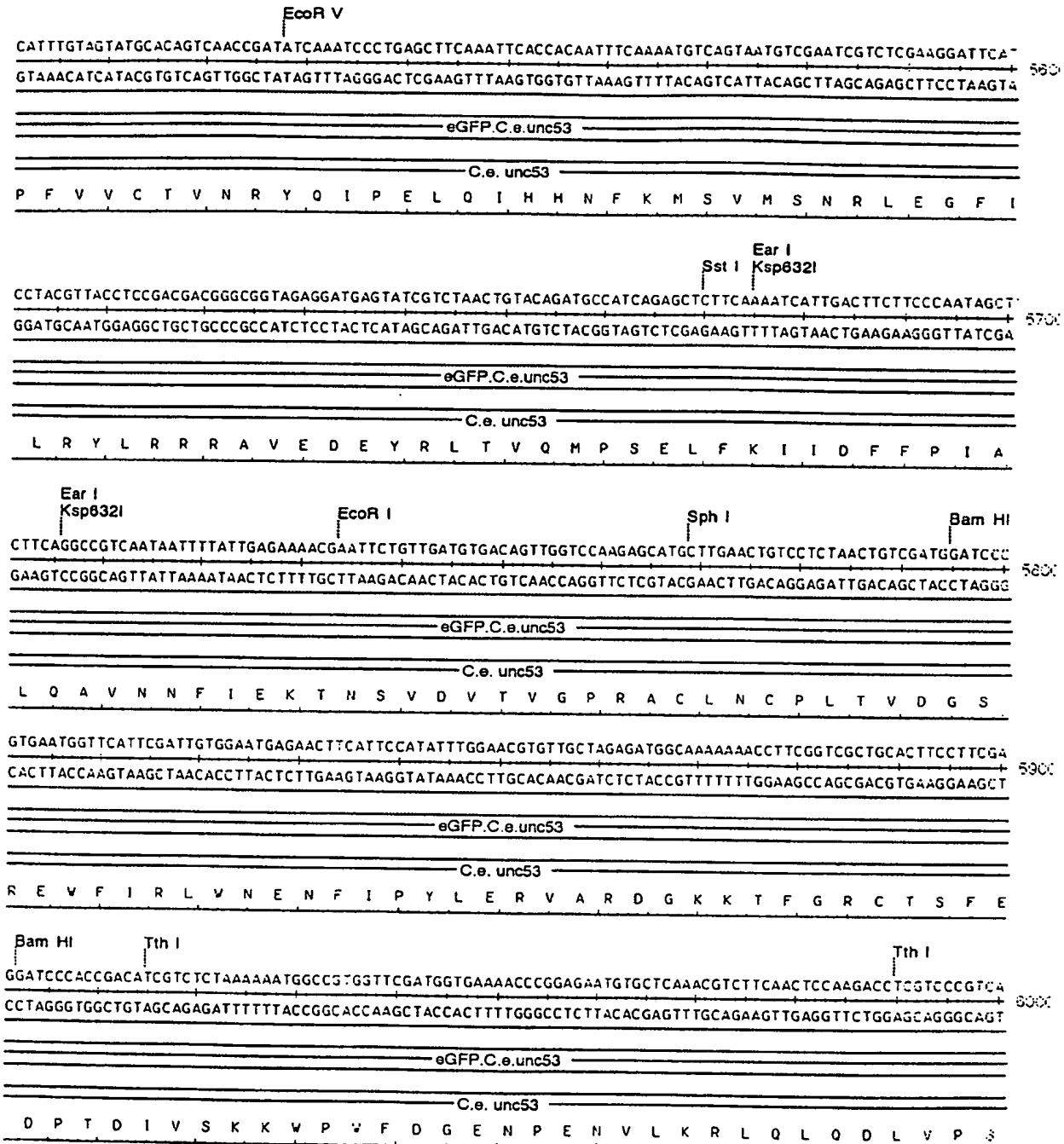
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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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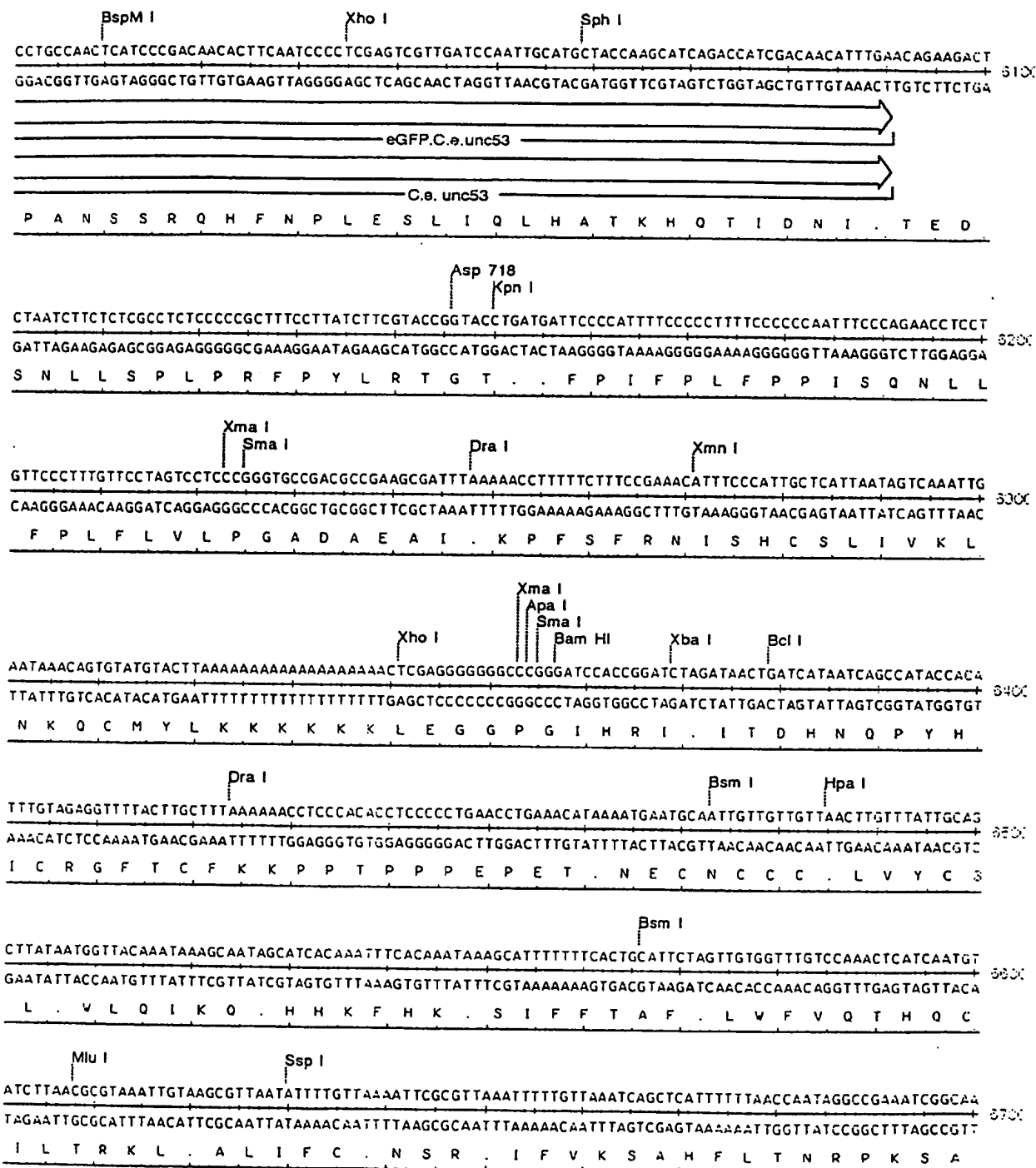


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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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Bsr I

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TTAGGGAATATTTAGTTTTCTTATCTGGCTCTATCCCAACTCACAACAAGSTCAAACCTTGTCTCAGGTGATAATTTCTTGACCTGAGGTTGCAGTTT
K S L I N Q K N R P R . G . V L F Q F G T R V H Y . R T V T P T S I 6800

Dra III

GGGCGAAAAACCGTCTATCAGGGCGATGGCCCACTACGTGAACCATCACCTAATCAAGTTTTTGGGGTCGAGGTGCCGTAAAGCACTAAATCGGAACC
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Nae I

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Ksp I

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Q V . R S R C A . P P H P P R L M R R Y R A R Q V A L F G E M C A E 7100

Bsp H I

Ssp I

Ear I
Ksp6321

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P L F V Y F S K Y I Q I C I R S . D N N P D K C F N N I E K G R V 7200

Oxa N I

Pvu II

Sph I
Ava III
Nsi I

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Sph I
Ava III
Nsi I

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Bsr I

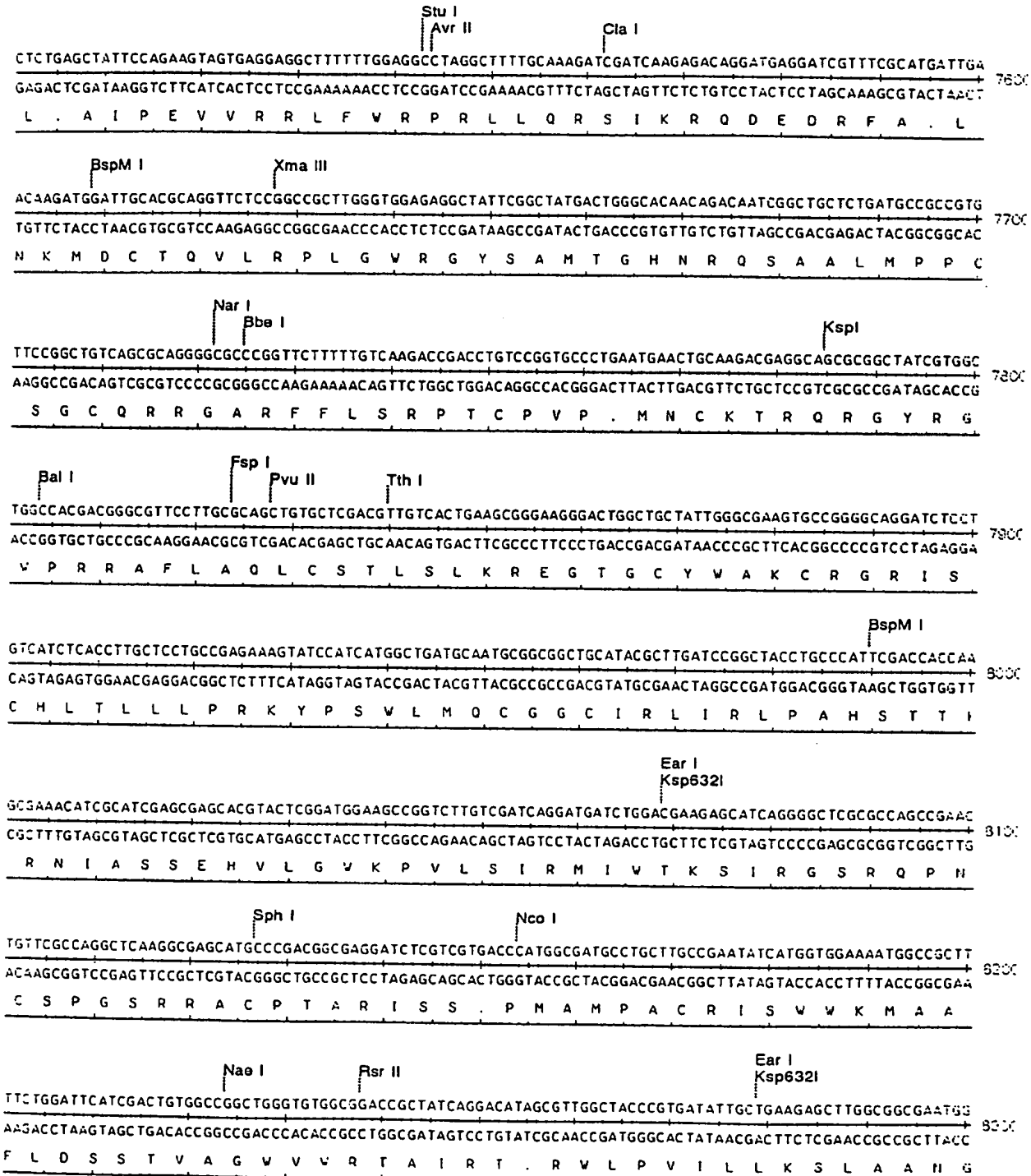
Nco I

Bgl I
Sfi I

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S A H P A P N S A Q F R P F S A P V L T N F F Y L C R G R G R L G 7500

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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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fig 30 pEGFP72 (1 > 9697) Site and Sequence

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GCTGACCGCTTCCTCGTGCTTTACGGTATCGCCGCTCCCGATTGCGAGCGCATCGCCTTCTATCGCCTTCTTGACGAGTTCTTCTGAGCGGGACTCTGSS
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Asu II BspM I
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V R N D R P S D A Q P A I T R F R F H R R L L . K V G L R N R F P

Nae I KspI Avr II
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G R R L D D P P A R G S H A G V L R P P . G E A N . N T E G D N T G

KspI
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R N P R Y D G N K K T E . N A R C V V V C S . T R G S V P G L A L

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C R Y P T E T P L G P I R P R F F L F P T P P P K F G . R P R A R

AlwI OxaN I Dra I Dra I
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S Q R R G G R P C H S L R L L I Y T L D . F K T S F L I . K D L G E

BspH I
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D P F . . S H D Q N P L T . V F V P L S V R P R R K D Q R I F L R

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BsrI
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E G N W L Q Q S A D T K Y C P S S V A V V R P P L Q E L C S T A Y I



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

| | | |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| (51) International Patent Classification ⁶ : C12N 15/12, 5/10, 15/85, C07K 14/435, 16/18, A61K 38/17, 49/00, C12Q 1/02, G01N 33/53 | A3 | (11) International Publication Number: WO 98/24810 (43) International Publication Date: 11 June 1998 (11.06.98) |
| (21) International Application Number: PCT/EP97/06956 (22) International Filing Date: 3 December 1997 (03.12.97) (30) Priority Data: 9625283.8 4 December 1996 (04.12.96) GB (71) Applicant (for all designated States except US): JANSSEN PHARMACEUTICA N.V. [BE/BE]; Turnhoutseweg 30, B-2340 Beerse (BE). (72) Inventors; and (75) Inventors/Applicants (for US only): PLATTEEUW, Christ, Jules [BE/BE]; Evergemsesteenweg 17, B-9032 Wondel- gem (BE). BUESA ARJOL, Carlos, Manuel [ES/ES]; Trav- essera de les Corts, 171/702a, E-08028 Barcelona (ES). DERAEMYAEKER, Marc [BE/BE]; Janssen Pharmaceu- tica N.V., Turnhoutseweg 30, B-2340 Beerse (BE). VER- HASSELT, Peter [BE/BE]; Janssen Pharmaceutica N.V., Turnhoutseweg 30, B-2340 Beerse (BE). PUJOL, Nathalie, Jeanne, Raymonde [FR/BE]; 213, avenue du Père Soulas, F-34000 Montpellier (FR). MAERTENS, Luc, Jacques, Si- mon [BE/BE]; Vier Uitersten 26, B-8200 Brugge (BE). LUYTEN, Walter [BE/BE]; Janssen Pharmaceutica N.V., Turnhoutseweg 30, B-2340 Beerse (BE). GEERTS, Hugo [BE/BE]; Janssen Pharmaceutica N.V., Turnhoutseweg 30, | | B-2340 Beerse (BE). VANDEKERCKHOVE, Joel, Ste- faan [BE/BE]; Rode Boukendreef 27, B-8210 Loppem (BE). GEYSEN, Johan [BE/BE]; Janssen Pharmaceutica N.V., Turnhoutseweg 30, B-2340 Beerse (BE). BOGAERT, Thierry, André, Olivier, Eddy [BE/BE]; Wolvendreef 26g, B-8500 Kortrijk (BE). (74) Agent: BALDOCK, Sharon, Claire; Boulton Wade Tennant, 27 Furnival Street, London EC4A 1PQ (GB). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims</i> <i>and to be republished in the event of the receipt of amendments.</i> (88) Date of publication of the international search report: 21 January 1999 (21.01.99) |
| (54) Title: VERTEBRATE HOMOLOGUES OF UNC-53 PROTEIN OF C. ELEGANS (57) Abstract Vertebrate protein homologues of UNC-53 protein of C. elegans and nucleic acid sequences coding for said homologues or functional equivalents thereof are identified. The nucleic acid sequences in an appropriate vector are used to transfect or transform cells, tissues or organisms useful in identifying inhibitors or enhancers of the vertebrate homologue, or further proteins involved in the signal transduction pathway of which said vertebrate homologue is a component. Any of said inhibitors or enhancers identified can be included in a pharmaceutical composition or in the preparation of a medicament for treating conditions such as neurological diseases, acute traumatic injuries and to promote neuronal regeneration and inhibit metastasis or loss of contact inhibition. | | |

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| DK | Denmark | LR | Liberia | SG | Singapore | | |
| EE | Estonia | | | | | | |

INTERNATIONAL SEARCH REPORT

Internat. Application No
PCT/EP 97/06956

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/12 C12N5/10 C12N15/85 C07K14/435 C07K16/18
A61K38/17 A61K49/00 C12Q1/02 G01N33/53

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|------------|----------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------|
| P, X | WO 96 38555 A (BOGAERT THIERRY (BE); STRINGHAM EVE (CA); VANDEKERCKHOVE JOEL (BE)) 5 December 1996 | 1, 2, 24-26, 28, 30-36, 38, 40, 42-51, 66, 68, 70, 72, 78, 79, 83-87, 98-105, 107, 112 |
| P, Y | see page 2, line 18 - page 20, line 26 | 3-23, 27, 29, 37, 39, 41, 58, 59, 67, 69, 74, 80-82, |
| -/-- | | |

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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- "&" document member of the same patent family

Date of the actual completion of the international search

2 October 1998

Date of mailing of the international search report

27. 11. 98

Name and mailing address of the ISA

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Fax: (+31-70) 340-3016

Authorized officer

Donath, C

INTERNATIONAL SEARCH REPORT

Internat'l Application No

PCT/EP 97/06956

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------|
| | <p>see figure 5</p> <p>see Sequence Listing: SEQ ID NO.: 1 and 4</p> <p>---</p> | <p>88, 115-118, 124,125</p> |
| Y | <p>HEKIMI, S. AND KERSHAW, D.: "Axonal guidance defects in a <i>Caenorhabditis elegans</i> mutant reveal cell-extrinsic determinants of neuronal morphology" THE JOURNAL OF NEUROSCIENCE, vol. 13, no. 10, October 1993, pages 4254-4271, XP000612286</p> <p>see page 4254, left-hand column, paragraph 1 - page 4255, right-hand column, paragraph 2</p> <p>see page 4267, right-hand column, paragraph 1 - page 4271, left-hand column, paragraph 2</p> <p>---</p> | <p>3-23,27, 29,37, 39,41, 58,59, 67,69, 74, 80-82, 88, 115-118, 124,125</p> |
| A | <p>STERN, M.J. ET AL.: "The human GRB2 and <i>Drosophila</i> Drk genes can functionally replace the <i>Caenorhabditis elegans</i> cell signaling gene <i>sem-5</i>" MOLECULAR BIOLOGY OF THE CELL, vol. 4, no. 11, November 1993, pages 1175-1188, XP002079466</p> <p>cited in the application</p> <p>see the whole document</p> <p>-----</p> | |

INTERNATIONAL SEARCH REPORT

International application No.
PCT/EP 97/06956

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Although claims 49-51, 58, 59, 107, 115 and 118 are directed to a diagnostic method practised on the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☒ Claims Nos.: 52-57, 60-65, 71, 73, 75-77, 89-94, 108-111, 113, 114
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Claims Nos.: 52-57,60-65,71,73,75-77,89-94,108-111,113,114

Claims 52-57, 60-65, 75-77, 91-94, 108-111,113 and 114 concern a compound, pharmaceutical composition, a nucleic acid sequence or the use of a compound. These compounds or compositions, however, are only defined by the method which can be used in order to identify these compounds or compositions as enhancers or inhibitors of regulation of cell shape, cell growth or motility or of the direction of cell migration or of the signal transduction pathway. Since it is completely unclear which kind of substances will be identified by the respective method and since in the specification no concrete examples for these kind of substances are given, the scope of said claims is totally ambiguous and undefined. Moreover, it cannot be excluded that even substances known in the art may be recognized as an enhancing or inhibiting compound by the respective used method.

The same applies to claims 71, 73, 89 and 90 concerning methods comprising proteins which are only defined by a reference to another method which was used to identify said proteins.

Information on patent family members

PCT/EP 97/06956

Form PCT/SA/210 (patent family annex) (July 1992)